

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: March 3, 2004, 12:11:47 ; Search time 11 Seconds

(without alignments)
23.668 Million cell updates/sec

Title: US-09-871-974-2

Perfect score: 29

Sequence: 1 TKPPR 5

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database : SwissProt_42.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	29	100.0	99	D127 HUMAN	Q9hlm4 homo sapien
2	29	100.0	201	COAE_SVNY3	Q5515 synchocyst
3	29	100.0	227	FA3C_HUMAN	Q95250 homo sapien
4	29	100.0	227	FA3C_MOUSE	Q91vu0 mus musculus
5	29	100.0	262	RPC_BPI63	P15238 bacterioph
6	29	100.0	465	ENGA_STRCO	Q9aww8 streptomyce
7	29	100.0	491	ENGA_STRAW	Q828y7 streptomyce
8	29	100.0	493	YE14 MYCPN	P75372 mycoplasma
9	29	100.0	604	YJ13 YEAST	P47030 saccharomyc
10	29	100.0	612	ELP1_MOUSE	Q60775 mus musculus
11	29	100.0	613	ELF1_HUMAN	P32519 homo sapien
12	29	100.0	919	DNLI_HUMAN	P18658 homo sapien
13	29	100.0	985	SEC8_DROME	Q9vnh6 drosophila
14	29	100.0	1067	BAB2_DROME	Q9w0k4 drosophila
15	29	100.0	1235	TRK1 YEAST	P13685 saccharomyc
16	29	100.0	1241	TRK1_SACBA	P28569 saccharomyc
17	26	89.7	54	RS14_PYRAE	Q8xvw1 pyrobaculum
18	26	89.7	133	SEC8_MOUSE	Q09535 mus musculus
19	26	89.7	133	VEGH_ORFN2	P52584 orf virus
20	26	89.7	140	RT12_DROME	P10735 drosophila
21	26	89.7	191	KGUA_SYNEL	Q8dmq7 synchococc
22	26	89.7	193	VC07_ADE04	Q96831 human adeno
23	26	89.7	198	VC07_ADE02	P03266 human adeno
24	26	89.7	199	KGUA_ANASP	Q82017 anabaena sp
25	26	89.7	205	KGUA_NEIMA	Q9jtc96 neisseria m
26	26	89.7	205	KGUA_NEIME	Q9jtp5 neisseria m
27	26	89.7	219	Y132_NPVAC	P24730 autocograph
28	26	89.7	220	PE55_LUCCU	Q95ue8 lucilia cup
29	26	89.7	253	NOCT_RAT	Q9et55 rattus norv
30	26	89.7	257	TRUA_XYLFA	Q9pdke xyella fas
31	26	89.7	257	TRUA_XYLFA	Q87ds1 xyella fas
32	26	89.7	304	YQOB_CAEEL	Q09300 caenorhabdi
33	26	89.7	350	YB04_AQUAE	Q67189 aquifex aeo

34	26	89.7	370	TRMU_RICCN	Q92i10 rickettsia
35	26	89.7	395	RT31_HUMAN	Q92665 homo sapien
36	26	89.7	396	TRMU_RHILO	Q98h10 rhizobium l
37	26	89.7	398	TRMU_BRUME	Q8yil6 brucella me
38	26	89.7	398	TRMU_BRUSU	Q8y38 brucella su
39	26	89.7	402	VGLD_HSVEA	P24872 equine herp
40	26	89.7	428	CCKR_HUMAN	P32238 homo sapien
41	26	89.7	429	NOCT_MOUSE	Q35710 mus musculu
42	26	89.7	431	NOCT_HUMAN	Q9uk39 homo sapien
43	26	89.7	438	ENGA_CLOPE	Q8xjk1 clostridium
44	26	89.7	439	SRMB_HAEIN	P44701 haemophilus
45	26	89.7	442	VGLD_HSVEK	P24484 equine herp
46	26	89.7	452	VGLD_HSVEB	P24379 equine herp
47	26	89.7	463	ENGA_BIFLO	Q8g6a8 bifidobacte
48	26	89.7	466	EN55_HUMAN	Q00013 homo sapien
49	26	89.7	484	ENP6_HUMAN	Q75354 homo sapien
50	26	89.7	524	ANPA_AESPE	Q9y935 aeropyrium p
51	26	89.7	536	GAG_MLVDB	P29168 murine leuk
52	26	89.7	552	AIRE_MOUSE	Q92063 mus musculu
53	26	89.7	619	OM70_NEUCR	P27961 neurospora
54	26	89.7	737	POLG_HCVJ5	P27960 hepatitis c
55	26	89.7	737	POLG_HCVJ7	P27961 hepatitis c
56	26	89.7	782	CS2B_HUMAN	Q9h8e8 homo sapien
57	26	89.7	846	IR81_HCVVA	P09715 human cytom
58	26	89.7	905	CIPF_MICTU	Q10860 mycobacteri
59	26	89.7	928	CHS2_EXODE	P36001 exophialad
60	26	89.7	961	LIN2_CAEEL	P54936 caenorhabdi
61	26	89.7	961	TSP4_HUMAN	P35443 homo sapien
62	26	89.7	991	DHP1_SCHPO	P40848 schizosacch
63	26	89.7	1057	POLR_DROME	P16423 drosophila
64	26	89.7	1122	VAB1_CAEEL	O61460 caenorhabdi
65	26	89.7	1130	ITR6_HUMAN	P32329 homo sapien
66	26	89.7	1160	WDR7_HUMAN	Q9y4e6 homo sapien
67	26	89.7	1173	DPOL_RCMYM	Q85428 rat cytoleg
68	26	89.7	1446	IE18_PVIFA	P33479 pseudorabie
69	26	89.7	1461	IE18_PVIFV	P11675 pseudorabie
70	26	89.7	2139	CCAC_MOUSE	Q01815 mus musculu
71	26	89.7	2169	CCAC_RAT	P22002 rattus norv
72	26	89.7	2171	CCAC_RABIT	P15381 oryctolagus
73	26	89.7	2221	CCAC_HUMAN	Q12936 homo sapien
74	26	89.7	2353	CCAH_HUMAN	Q95180 homo sapien
75	26	89.7	3033	POLG_HCVJ8	P26661 h genome po
76	26	89.7	5262	MLI2_HUMAN	O14686 homo sapien
77	25	86.2	44	COXR_RAT	P80433 rattus norv
78	25	86.2	69	COXR_BOVIN	P14622 bos taurus
79	25	86.2	111	LV21_HUMAN	P01712 homo sapien
80	25	86.2	114	MAUL_METEX	Q9129 methylobact
81	25	86.2	172	RUB2_PSEOL	P00272 pseudomonas
82	25	86.2	223	GLU2_MAIZE	P04706 zea mays (m
83	25	86.2	228	MX11_HUMAN	P50539 homo sapien
84	25	86.2	228	MX11_MOUSE	P50540 mus musculu
85	25	86.2	228	MX11_RAT	O09015 rattus norv
86	25	86.2	241	NGF_HUMAN	P01138 homo sapien
87	25	86.2	262	NEF_SIVM1	P05862 simian immu
88	25	86.2	277	TRMD_STRCO	O69882 streptomyce
89	25	86.2	278	PYRF_BACSL	Q8nq40 corynebacte
90	25	86.2	296	YWFH_BACSU	P39649 bacillus su
91	25	86.2	297	SGS4_DROME	Q00725 drosophila
92	25	86.2	306	FMT_BRUME	Q8ydb3 brucella me
93	25	86.2	311	FMT_AGR75	Q8uid0 agrobacteri
94	25	86.2	311	FMT_RHIME	Q92sh5 rhizobium m
95	25	86.2	317	FMT_RHILO	Q98d53 rhizobium l
96	25	86.2	318	IFR_CICAR	Q00016 cicier ariet
97	25	86.2	332	SR4_PHYPO	P11113 physarum po
98	25	86.2	332	YIE2_HSVB4	Q02484 bovine herp
99	25	86.2	379	GIT7_SCHPO	O59709 schizosacch
100	25	86.2	396	PSPB_DICDI	P54704 dictyosteli
101	25	86.2	396	TRT_DROME	P19351 drosophila
102	25	86.2	412	SUS1_MOUSE	O54864 m histone-1
103	25	86.2	431	ACRO_RABIT	P48038 oryctolagus
104	25	86.2	434	UL43_HSV11	P10227 herpes simp
105	25	86.2	458	ENGA_HELPY	O25505 helicobacte
106	25	86.2	462	ENGA_HELPY	Q92109 helicobacte

```

107 25 86.2 469 1 ANTA_GENTR
108 25 86.2 487 1 CPDE_BOVIN
109 25 86.2 499 1 DCE_CANFA
110 25 86.2 502 1 DCE_LYCES
111 25 86.2 607 1 YK06_CABEL
112 25 86.2 649 1 TOP3_VISCH
113 25 86.2 649 1 TOP3_SALTY
114 25 86.2 649 1 TOP3_SALTY
115 25 86.2 653 1 TOP3_ECOLI
116 25 86.2 672 1 NKR1_YEAST
117 25 86.2 679 1 ENV2_MOUSE
118 25 86.2 908 1 SRCA_RABIT
119 25 86.2 988 1 ST23_YEAST
120 25 86.2 1035 1 C68_YEAST
121 25 86.2 1074 1 EM12_MOUSE
122 25 86.2 1080 1 NKR1_YEAST
123 25 86.2 1152 1 ITAM_HUMAN
124 25 86.2 1175 1 DSRA_FAT
125 25 86.2 1210 1 AP4_HUMAN
126 25 86.2 1276 1 BMD_CLOBO
127 25 86.2 1411 1 TCOF_HUMAN
128 25 86.2 1484 1 CES2_HUMAN
129 25 86.2 1556 1 Y934_HUMAN
130 25 86.2 2146 1 INSR_DROME
131 25 86.2 2472 1 NCR2_MOUSE
132 25 86.2 5596 1 WDR1_HUMAN
133 24 82.8 57 1 Y23A_MYCLE
134 24 82.8 59 1 MAMB_DENJA
135 24 82.8 60 1 RZ0D_ECOLI
136 24 82.8 61 1 TX51_DENJA
137 24 82.8 62 1 ITR1_ASCSU
138 24 82.8 63 1 YAL1_ECOLI
139 24 82.8 67 1 SLX1_RHILLO
140 24 82.8 68 1 SLX1_BRUME
141 24 82.8 68 1 SLX1_BRUSU
142 24 82.8 73 1 UL12_HCMVA
143 24 82.8 80 1 SSS2_SCYCA
144 24 82.8 91 1 RL31_PYRAE
145 24 82.8 93 1 N12A_MEDSA
146 24 82.8 96 1 YBCO_ECOLI
147 24 82.8 103 1 N012_MEDTR
148 24 82.8 105 1 THGF_TOBAC
149 24 82.8 106 1 THG1_NICPA
150 24 82.8 107 1 REX1_SCHPO

ALIGNMENTS
ID D127_HUMAN STANDARD; PRT; 99 AA.
AC Q9HLM4;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DE Beta-defensin 127 precursor (Beta-defensin 27) (DEFB-27).
GN DEFB127 OR DEFB27 OR C20ORF73.
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP MEDLINE=21638749; PubMed=11780052;
RA Deloukas P., Matthews L.H., Ashurst J., Burton J., Gilbert J.G.R.,
RA Jones M., Stavrides G., Almeida J.P., Babbage A.K., Begguley C.L.,
RA Bailey J., Barlow K.F., Bates K.N., Beard L.M., Beare D.M.,
RA Beasley O.P., Bird C.P., Blakey S.E., Bridgeman A.M., Brown A.J.,
RA Buck D., Burrill W.D., Butler A.P., Carder C., Carter N.P.,
RA Chapman J.C., Clamp M., Clark G., Clark L.N., Clark S.Y., Clee C.M.,
RA Clegg S., Colley V.E., Collier R.E., Connor R.E., Corby N.R.,
RA Coulson A., Coville G.J., Deadman R., Dhani P.D., Dunn M.,

```

Ellington A.G., Frankland J.A., Fraser A., French L., Garner P.,
Graham D.V., Griffiths C., Griffiths M.N.D., Gwilliam R., Hall R.E.,
Hammond S., Hartley J.L., Heath P.D., Ho S., Holden J.L., Howden P.J.,
Huckle E., Hunt A.R., Hunt S.B., Jekosch K., Johnson C.M., Johnson D.,
Kay M.P., Kimberley A.M., King A., Knights A., Laird G.K., Lawlor S.,
Lehvaeslahti M.H., Leversha M.A., Lloyd C., Lloyd D.M., Lovell J.D.,
Marsh V.L., Martin S.L., McConnachie I.J., McElay K., McMurray A.A.,
Milne S.A., Mistry D., Moore W.J.P., Mullikin J.C., Nickerson T.,
Oliver K., Parker A., Patel R., Pearce T.A.V., Peck A.I.,
Phillimore B.U.C.T., Prathalingam S.R., Plumb R.W., Ramsay H.,
Rice C.M., Ross M.T., Scott C.E., Sehra H.K., Showkhen R., Sims S.,
Skuce C.D., Smith M.L., Soderlund C., Steward C.A., Sulston J.E.,
Swann R.M., Sycamore N., Taylor R., Tee L., Thomas D.W., Thorpe A.,
Tracey A., Tromans A.C., Vaudin M., Wall M., Wallis J.M.,
Whitehead S.L., Whitaker P., Willey D.L., Williams L., Williams S.A.,
Wilming L., Wray P.W., Hubbard T., Durbin R.M., Bentley D.R., Beck S.,
Rogers J.,
The DNA sequence and comparative analysis of human chromosome 20.";
Nature 414:865-871 (2001).

SEQUENCE OF 19-77 FROM N.A., AND IDENTIFICATION.
TISSUE=B-cell, Petal lung, and Testis;
MEDLINE=21843921; PubMed=11854508;
Schutte B.C., Mitros J.P., Bartlett J.A., Walters J.D., Jia H.P.,
Welsh M.J., Casavant T.L., McCray P.B. Jr.;
"Discovery of five conserved beta-defensin gene clusters using a
computational search strategy.";
Proc. Natl. Acad. Sci. U.S.A. 99:2129-2133 (2002).
CC -!- FUNCTION: Has antibacterial activity (potential).
CC -!- SUBCELLULAR LOCATION: Secreted (potential).
CC -!- SIMILARITY: Belongs to the beta-defensin family.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
between the Swiss Institute of Bioinformatics and the EMBL outstation -
the European Bioinformatics Institute. There are no restrictions on its
use by non-profit institutions as long as its content is in no way
modified and this statement is not removed. Usage by and for commercial
entities requires a license agreement (see http://www.isb-sib.ch/announce/
or send an email to license@isb-sib.ch).
EMBL; AL360078; CACU7685.1;
EMBL; AY122479; AAM3930.1;
Genew; HGNC:16206; DEFB127.
DR GO; GO:0003797; F:antibacterial peptide activity; TAS.
DR GO; GO:0045087; P:innate immune response; TAS.
KW Antibiotic; Signal.
FT SIGNAL 1
FT CHAIN 19 63
FT PROPEP 66 99
FT DISULFID 24 53
FT DISULFID 33 47
FT DISULFID 37 54
FT CONFLICT 31 31
FT CONFLICT 71 77
SQ SEQUENCE 99 AA; 11342 MW; C9CDB27D17AF380 CRC64;
Query Match 100.0%; Score 29; DB 1; Length 99;
Best Local Similarity 100.0%; Pred. No. 22; Indels 0; Gaps 0;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
Db 67 TKPPR 71
RESULT 2
ID COAE_SYNY3 STANDARD; PRT; 201 AA.
AC Q55515;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Dephospho-CoA kinase (EC 2.7.1.24) (Dephosphocoenzyme A kinase).

GN COAE OR SLR0553.
 OS Synechocystis sp. (strain PCC 6803).
 OC Bacteria; Cyanobacteria; Chroococcales; Synechocystis.
 CX NCBI_TaxID=1148;
 RX SEQUENCE FROM N.A.
 RP MEDLINE=96127429; PubMed=8590279;
 RA Kaneko T., Tanaka A., Sato S., Kotani H., Sazuka T., Miyajima N.,
 RT "Sequence analysis of the genome of the unicellular cyanobacterium
 RT Synechocystis sp. strain PCC6803. I. Sequence features in the 1 Mb
 RT region from map positions 64% to 92% of the genome.";
 RL DNA Res. 2:153-166(1995).
 CC -1- FUNCTION: Catalyzes the phosphorylation of the 3'-hydroxyl group
 CC of dephosphocoenzyme A to form coenzyme A (By similarity).
 CC -1- CATALYTIC ACTIVITY: ATP + dephospho-CoA = ADP + CoA.
 CC -1- PATHWAY: Coenzyme A (CoA) biosynthesis; Fifth (last) step.
 CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
 CC -1- SIMILARITY: Belongs to the coae family.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 DR EMBL: D64006; BAAL0873.1; -.
 DR HAMAP: MF_00376; -; 1.
 DR InterPro: IPR001977; Depp_CoKinase.
 DR Pfam: PF01121; Coae; 1. CoKinase; 1.
 DR PRODOM: PD003329; Depp_CoKinase; 1.
 DR TIGRGRAMS: TIGR00152; TIGR00152; 1.
 DR PROSITE: PS01294; COAE; 1.
 DR Transferrase; Kinase; ATP-binding; Coenzyme A biosynthesis;
 KW Complete proteome.
 FT NP_BIND 15 22 ATP (POTENTIAL).
 SQ SEQUENCE 201 AA; 22520 MW; 34C3E142337F519E CRC64;
 Query Match 100.0%; Score 29; DB 1; Length 201;
 Best Local Similarity 100.0%; Pred. No. 45;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPR 5
 DB 3 TKPR 7
 RESULT 3
 FA3C HUMAN STANDARD; PRT; 227 AA.
 AC Q92520;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Protein FAM3C precursor (Protein GS3786).
 GN FAM3C.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 CX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Bone;
 RA Ono I., Hashimoto J., Takaoka K., Ochi T., Okubo K., Matsubara K.;
 RL Submitted (AUG-1996) to the EMBL/GenBank/DBJ databases.
 [2]
 RP SEQUENCE FROM N.A.
 RA Ryan E., Bauer C., Tucci S., Spalding L.;
 RL Submitted (JAN-1999) to the EMBL/GenBank/DBJ databases.
 [3]

RP SEQUENCE FROM N.A.
 RC TISSUE=Ductum;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Krausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Colling F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Bustow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Datchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalón D.K., Murthy D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Kettner M., Madan A., Rodriguez S., Sanchez A.,
 RA Whitney M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smallos D.E.,
 RA Schnurch A., Schein J.E., Jones S.J.M., Marra M.A.,
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 CC CHARACTERIZATION.
 CC MEDLINE=22150857; PubMed=12160727;
 RA Zhu Y., Xu G., Patel A., McLaughlin M.M., Silverman C., Knecht K.A.,
 RA Sweitzer S., Li X., McDonnell P., Mirabile R., Zimmerman D., Boyce R.,
 RA Tierney L.A., Hu E., Livi G.P., Wolf B.A., Abdel-Meguid S.S.,
 RA Rose G.D., Aurora R., Hensley P., Briggs M., Young P.R.,
 RT "Cloning, expression, and initial characterization of a novel
 RT cytokine-like gene family.";
 RL Genomics 80:144-150(2002).
 CC -1- SUBCELLULAR LOCATION: Secreted (Potential).
 CC -1- TISSUE SPECIFICITY: Ubiquitous.
 CC -1- SIMILARITY: Belongs to the FAM3 family.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 DR EMBL: D87120; BAAL3251.1; -.
 DR EMBL: AC006364; AAQ96872.1; -.
 DR EMBL: BC046932; AAF46932.1; -.
 DR Genew; HGNC:18664; FAM3C.
 DR GO: GO:0005576; C:extracellular; NAS.
 DR GO: GO:0005125; F:cytokine activity; NAS.
 KW Signal.
 FT SIGNAL 1 24 POTENTIAL.
 FT CHAIN 25 227 PROTEIN FAM3C.
 FT DISULFID 58 221 POTENTIAL.
 FT DISULFID 64 86 POTENTIAL.
 SQ SEQUENCE 227 AA; 24680 MW; 6DC94B259052647F CRC64;
 Query Match 100.0%; Score 29; DB 1; Length 227;
 Best Local Similarity 100.0%; Pred. No. 50;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPR 5
 DB 51 TKPR 55
 RESULT 4
 FA3C MOUSE STANDARD; PRT; 227 AA.
 ID FA3C MOUSE
 AC Q91VU0; Q9CTB4;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)

```

DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Protein FAM3C precursor.
GN FAM3C OR D6WSU176E.
OS Mus musculus (Mouse).
CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
CX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain tumor;
PX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Haie H.,
RA Diatchenko L., Marusina K., Fawcett A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldi M.P., Casavant T.L., Scheetz T.E.,
RA Raba S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaby S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Whiting M., Madan A., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.J., Skalska U., Smailus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RA "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP SEQUENCE OF 127-227 FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Embryo;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamataka I.,
RA Saito T., Okazaki Y., Gotojori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
RA Fleischmann W., Gaasterland T., Glass C., King B., Kochiwa H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
RA Schirni L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Sakamoto N.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whitaker C., Wilming L.,
RA Wyszynski-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:695-690(2001).
CC -!- SUBCELLULAR LOCATION: Secreted (Potential).
CC -!- SIMILARITY: Belongs to the FAM3 family.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; BC009086.1; -
DR EMBL; AK004059; BAB23146.1; -
DR MGD; MGI:107392; D6WSU176E;
DR GO; GO:0005576; C:extracellular; ISS.
DR GO; GO:0005125; F:cytokine activity; ISS.
FT SIGNAL.
1 24 POTENTIAL.

FT CHAIN 25 227 PROTEIN FAM3C.
FT DISULFID 58 221 POTENTIAL.
FT DISULFID 64 86 POTENTIAL.
SQ SEQUENCE 227 AA; 24752 MW; 9195280B92838CF4 CRC64;

Query Match 100.0%; Score 29; DB 1; Length 227;
Best Local Similarity 100.0%; Pred. No. 50;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 51 TKPPR 55

RESULT 5
ID RPC BP163 STANDARD; PRT; 262 AA.
AC P15238; O9MCD02;
DT 01-APR-1990 (Rel. 14, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Repressor protein C.
GN C.
OS Bacteriophage 16-3.
OC Viruses; dsDNA viruses, no RNA stage; Caudovirales.
OX NCBI_TaxID=10704;
RN [1]
RP SEQUENCE FROM N.A., AND SEQUENCE OF 1-3.
RA Dallmann G., Papp P., Oroz L.;
RT "Related repressor specificity of unrelated phages.";
RL Nature 330:398-401(1987).
RN [2]
RP REVISIONS.
RX MEDLINE=99328962; PubMed=10400574;
RA Seney S., Papp I., Buzas Z., Papp P.;
RT "Identification of site-specific recombination genes int and xis of
RT the Rhizobium temperate phage 16-3.";
RL J. Bacteriol. 181:4185-4192(1999).
CC -!- SIMILARITY: Contains 1 HTH cro/C1-type DNA-binding domain.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; AJ131679; CAB54833.1; -
DR PIR; S01612; RPB216.
DR InterPro; IPR001387; HTH_3.
DR Pfam; PF01381; HTH_3; 1.
DR SMART; SM00530; HTH_XRE; 1.
DR PROSITE; PS00943; HTH_CROCI; 1.
DR Early protein; Transcription regulation; Repressor; DNA-binding.
FW INIT MET 0 0
FT DOMAIN 14 67 HTH CRO/C1-TYPE.
FT DNA BIND 25 44 H-T-H MOTIF (PROBABLE).
SQ SEQUENCE 262 AA; 29522 MW; 56DE1FF97AC9C010 CRC64;

Query Match 100.0%; Score 29; DB 1; Length 262;
Best Local Similarity 100.0%; Pred. No. 58;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 192 TKPPR 196

RESULT 6
ID ENGA STRCO STANDARD; PRT; 465 AA.
AC Q9EW8;

```

28-FEB-2003 (Rel. 41, Created)
 28-FEB-2003 (Rel. 41, Last sequence update)
 15-MAR-2004 (Rel. 43, Last annotation update)
 GTP-binding protein enga.
 ENGA OR SC01758 OR 28CI34.11C.
 Streptomyces coelicolor.
 Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
 Streptomycineae; Streptomycetaceae; Streptomyces.
 NCBI_TaxID=1902;
 [1]
 SEQUENCE FROM N.A.
 STRAIN=AS(2) / M145;
 MEDLINE=21996410; PubMed=12000953;
 Bentley S.D., Chater K.F., Cerdano-Tarraga A.-M., Challis G.L.,
 Thomson N.R., James K.D., Harris D.E., Quail M.A., Kieser H.,
 Harper D., Bateman A., Brown S., Chandra G., Chen C.W., Collins M.,
 Cronin A., Fraser A., Goble A., Hidalgo J., Hornsby T., Howarth S.,
 Huang C.-H., Kieser T., Larke L., Murphy L., Oliver K., O'Neill S.,
 Rabinowitsch E., Rajandream M.A., Rutherford K., Rutter S.,
 Seeger K., Saunders D., Sharp S., Squares R., Squares S., Taylor K.,
 Warren T., Wietzorrek A., Woodward J., Barrell B.G., Parkhill J.,
 Hopwood D.A.;
 "Complete genome sequence of the model actinomycete Streptomyces
 coelicolor A3(2).";
 Nature 417:141-147(2002).
 -!- FUNCTION: GTPase of unknown physiological role.
 -!- SIMILARITY: Belongs to the era/tme family of GTP-binding
 proteins. Enga subfamily.

 This SWISS-PROT entry is copyright. It is produced through a collaboration
 between the Swiss Institute of Bioinformatics and the EMBL outstation
 at the European Bioinformatics Institute. There are no restrictions on its
 use by non-profit institutions as long as its content is in no way
 modified and this statement is not removed. Usage by and for commercial
 entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 or send an email to license@isb-sib.ch).

 EMBL: AL393110; CAC12931.1; ALT INIT.
 HAMAP: MF 00195; -; 1.
 InterPro: IPR003593; AAA ATPase.
 InterPro: IPR005289; GTP-binding_dom.
 InterPro: IPR006073; GTP1_OBG.
 InterPro: IPR002917; MMR_HSR1.
 Pfam: PF01926; MMR_HSR1; 1.
 PRINTS: PR00326; GTP1_OBG.
 SMART: SM00382; AAA; 2.
 TIGRFAMs: TIGR00650; MG442; 2.
 TIGRFAMs: TIGR00231; small_GTP; 2.
 GTP-binding; Repeat; Complete proteome.
 NP_BIND 33 40 GTP 1 (POTENTIAL).
 FT NP_BIND 80 84 GTP 1 (POTENTIAL).
 FT NP_BIND 142 145 GTP 1 (POTENTIAL).
 FT NP_BIND 208 215 GTP 2 (POTENTIAL).
 FT NP_BIND 255 259 GTP 2 (POTENTIAL).
 FT NP_BIND 320 323 GTP 2 (POTENTIAL).
 FT NP_BIND 465 465 GTP 2 (POTENTIAL).
 SQ SEQUENCE 465 AA; 50344 MW; 95053E7C00C859F8 CRC64;
 Query Match 100.0%; Score 29; DB 1; Length 465;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 |
 |
 |
 |
 DB 414 TKPPR 418
 RESULT 7
 ENGA_STRAW
 ID ENGA_STRAW STANDARD; PRT; 491 AA.
 AC Q828Y7;
 DT 15-MAR-2004 (Rel. 43, Created)
 DT 15-MAR-2004 (Rel. 43, Last sequence update)

15-MAR-2004 (Rel. 43, Last annotation update)
 GTP-binding protein enga.
 ENGA OR SAV6524.
 Streptomyces avermitilis.
 Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
 Streptomycineae; Streptomycetaceae; Streptomyces.
 NCBI_TaxID=33903;
 [1]
 SEQUENCE FROM N.A.
 STRAIN=NA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
 MEDLINE=21477403; PubMed=11572948;
 Omura S., Ikeda H., Ishikawa J., Hanamoto A., Takahashi C.,
 Shinose M., Takahashi Y., Horikawa H., Nakazawa H., Osonoe T.,
 Kikuchi H., Shiba T., Sakaki Y., Hattori M.;
 "Genome sequence of an industrial microorganism Streptomyces
 avermitilis: deducing the ability of producing Streptomyces
 metabolites.";
 Proc. Natl. Acad. Sci. U.S.A. 98:12215-12220(2001).
 [2]
 SEQUENCE FROM N.A.
 STRAIN=NA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
 MEDLINE=22608306; PubMed=13692562;
 Ikeda H., Ishikawa J., Hanamoto A., Shinose M., Kikuchi H., Shiba T.,
 Sakaki Y., Hattori M., Omura S.;
 "Complete genome sequence and comparative analysis of the industrial
 microorganism Streptomyces avermitilis.";
 Nat. Biotechnol. 21:526-531(2003).
 -!- FUNCTION: GTPase of unknown physiological role.
 -!- SIMILARITY: Belongs to the era/tme family of GTP-binding
 proteins. Enga subfamily.

 This SWISS-PROT entry is copyright. It is produced through a collaboration
 between the Swiss Institute of Bioinformatics and the EMBL outstation
 at the European Bioinformatics Institute. There are no restrictions on its
 use by non-profit institutions as long as its content is in no way
 modified and this statement is not removed. Usage by and for commercial
 entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 or send an email to license@isb-sib.ch).

 EMBL: AP005047; BAC74235.1; -.
 HAMAP: MF 00195; -; 1.
 InterPro: IPR003593; AAA ATPase.
 InterPro: IPR005289; GTP-binding_dom.
 InterPro: IPR006073; GTP1_OBG.
 InterPro: IPR002917; MMR_HSR1.
 Pfam: PF01926; MMR_HSR1; 1.
 PRINTS: PR00326; GTP1_OBG.
 SMART: SM00382; AAA; 2.
 TIGRFAMs: TIGR00650; MG442; 2.
 TIGRFAMs: TIGR00231; small_GTP; 2.
 GTP-binding; Repeat; Complete proteome.
 NP_BIND 59 66 GTP 1 (POTENTIAL).
 FT NP_BIND 106 110 GTP 1 (POTENTIAL).
 FT NP_BIND 168 171 GTP 1 (POTENTIAL).
 FT NP_BIND 234 241 GTP 2 (POTENTIAL).
 FT NP_BIND 281 285 GTP 2 (POTENTIAL).
 FT NP_BIND 346 349 GTP 2 (POTENTIAL).
 SQ SEQUENCE 491 AA; 53353 MW; 7BADD8E0449D14AC CRC64;
 Query Match 100.0%; Score 29; DB 1; Length 491;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 |
 |
 |
 |
 DB 440 TKPPR 444
 RESULT 8
 YE14_MYCPN
 ID YE14_MYCPN STANDARD; PRT; 493 AA.
 AC P75372;

DR InterPro; IPR000418; ETS.
 DR InterPro; IPR002341; HSF.ETS.
 DR Pfam; PF00178; Ets; 1.
 DR PRINTS; PR00454; ETSDOMAIN.
 DR SMART; SM00413; ETS; 1.
 DR PROSITE; PS00345; ETS DOMAIN 1; 1.
 DR PROSITE; PS00346; ETS DOMAIN 2; 1.
 DR PROSITE; PS0061; ETS DOMAIN 3; 1.
 DR Nuclear protein; Transcription regulation; Activator; DNA-binding.
 KW DOMAIN 75 80 POLY-ASP.
 FT DNA BIND 208 290
 FT SEQUENCE 612 AA; 66221 MW; 442F4C95142B31F0 CRC64;
 SQ

Query Match 100.0%; Score 29; DB 1; Length 612;
 Best Local Similarity 100.0%; Pred. No. 1.4e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
 Db 180 TKPPR 184

RESULT 11
 ETL1 HUMAN
 ID ETL1 HUMAN STANDARD; PRT; 619 AA.
 AC P32519; Q9UDE1;
 DT 01-OCT-1993 (Rel. 27, Created)
 DT 01-OCT-1993 (Rel. 27, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE ETS-related transcription factor Etl-1 (E74-like factor 1).
 GN ETL1.
 OS Homo sapiens (Human).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=92407982; PubMed=1527846;
 RA Leiden J.M., Wang C.Y., Petryniak B., Markovitz D.M., Nabel G.J.,
 RA Thompson C.B.;
 RA "A novel Ets-related transcription factor, Etl-1, binds to human
 RT immunodeficiency virus type 2 regulatory elements that are required
 RT for inducible trans activation in T cells.";
 RL J. Virol. 66:5890-5897(1992).
 RN [2]
 RP SEQUENCE OF 204-289 FROM N.A.
 RX MEDLINE=92186836; PubMed=1545787;
 RA Thompson C.B., Wang C.Y., Ho I.C., Bohjanen P.R., Petryniak B.,
 RA June C.H., Miesfeldt S., Zhang L., Nabel G.J., Karpinski B.;
 RT "Cis-acting sequences required for inducible interleukin-2 enhancer
 RT function bind a novel Ets-related protein, Etl-1.";
 RL Mol. Cell. Biol. 12:1043-1053(1992).
 RN [3]
 RP BINDING TO RB.
 RX MEDLINE=93262492; PubMed=8493578;
 RA Wang C.Y., Petryniak B., Thompson C.B., Kaelin W.G., Leiden J.M.;
 RT "Regulation of the Ets-related transcription factor Etl-1 by binding
 RT to the retinoblastoma protein.";
 RL Science 260:1330-1335(1993).
 CC -!- FUNCTION: TRANSCRIPTION FACTOR THAT APPEARS TO BE REQUIRED FOR THE
 CC T-CELL-RECEPTOR-MEDIATED TRANS ACTIVATION OF HIV-2 GENE
 CC EXPRESSION. BINDS SPECIFICALLY TO TWO PURINE-RICH MOTIFS IN THE
 CC HIV-2 ENHANCER. ETL-1 BINDS TO THE UNDERPHOSPHORYLATED FORM OF RB.
 CC MAY INTERACT WITH OTHER TRANSCRIPTION FACTORS IN ORDER TO REGULATE
 CC SPECIFIC GENES.
 CC -!- SUBCELLULAR LOCATION: Nuclear.
 CC -!- SIMILARITY: Belongs to the ETS family.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial

CC entities requires a license agreement (see http://www.isb-sib.ch/announce/
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; M82882; -; NOT_ANNOTATED_CDS.
 CC FIr; A43361; A43361.
 DR HSP; P28324; IBC8.
 DR TRANSFAC; T01113; -.
 DR Genew; HGNC:3316; ETL1.
 DR MTM; 189973; -.
 DR GO; GO:0005634; C:nucleus; NAS.
 DR GO; GO:0015663; F:transcriptional activator activity; NAS.
 DR GO; GO:0006355; P:regulation of transcription, DNA-dependent; NAS.
 DR InterPro; IPR000418; HSF.ETS.
 DR InterPro; IPR002341; HSF.ETS.
 DR Pfam; PF00178; Ets; 1.
 DR PRINTS; PR00454; ETSDOMAIN.
 DR SMART; SM00413; ETS; 1.
 DR PROSITE; PS00345; ETS DOMAIN 1; 1.
 DR PROSITE; PS00346; ETS DOMAIN 2; 1.
 DR PROSITE; PS0061; ETS DOMAIN 3; 1.
 KW Nuclear protein; Transcription regulation; Activator; DNA-binding.
 FT DOMAIN 75 80 POLY-ASP.
 FT DNA BIND 208 290
 FT CONFLICT 283 283 Q -> G (IN REF. 2).
 FT SEQUENCE 619 AA; 67455 MW; AB0B41B2964A66EF CRC64;
 SQ

Query Match 100.0%; Score 29; DB 1; Length 619;
 Best Local Similarity 100.0%; Pred. No. 1.4e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
 Db 180 TKPPR 184

RESULT 12
 DNLI HUMAN
 ID DNLI HUMAN STANDARD; PRT; 919 AA.
 AC P18858;
 DT 01-NOV-1990 (Rel. 16, Created)
 DT 01-NOV-1990 (Rel. 16, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE DNA ligase I (EC 6.5.1.1) (Polydeoxyribonucleotide synthase [ATP]).
 GN LIG1.
 OS Homo sapiens (Human).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=T lymphoblast;
 RX MEDLINE=90370849; PubMed=2204063;
 RA Barnes D.E., Johnston L.H., Kodama K.I., Tomkinson A.E.,
 RA Lasko D.D., Lindahl T.;
 RA "Human DNA ligase I cDNA: cloning and functional expression in
 RT Saccharomyces cerevisiae.";
 RL Proc. Natl. Acad. Sci. U.S.A. 87:6679-6683(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Rieder M.J., Livingston R.J., Braun A.C., Montoya M.A., Chung M.-W.,
 RA Miyamoto K.E., Nguyen C.P., Nguyen D.A., Poel C.L., Robertson P.D.,
 RA Schackwitz W.S., Sherwood J.K., Witrak L.A., Nickerson D.A.;
 RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE OF 716-753 FROM N.A.
 RX MEDLINE=91352039; PubMed=1881902;
 RA Petrin J.H.J., Huwiler K.G., Weaver D.T.;
 RA "A wild-type DNA ligase I gene is expressed in Bloom's syndrome
 RT cells.";
 RL Proc. Natl. Acad. Sci. U.S.A. 88:7615-7619(1991).
 RN [4]
 RP VARIANTS LYS-566 AND TRP-771.
 RX MEDLINE=92257590; PubMed=1581963;

RA Barnes D.E., Tomkinson A.E., Lehmann A.R., Webster A.D.B.,
 RA Lindahl T.;
 RT "Mutations in the DNA ligase I gene of an individual with
 RT immunodeficiencies and cellular hypersensitivity to DNA-damaging
 RL agents";
 CC Cell 69:495-503(1992).
 CC -!- FUNCTION: This protein seals, during DNA replication, DNA
 CC recombination and DNA repair, nicks in double-stranded DNA.
 CC -!- CATALYTIC ACTIVITY: ATP + (deoxyribonucleotide)(N) +
 CC (deoxyribonucleotide)(N-M)
 CC -!- SUBCELLULAR LOCATION: Nuclear.
 CC -!- PM: Phosphorylated in vivo.
 CC -!- DISEASE: Defects in LIG1 seem to cause immunodeficiencies and
 CC cellular hypersensitivity to DNA-damaging agents.
 CC -!- SIMILARITY: Belongs to the ATP-dependent DNA ligase family.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; M36067; AAA59518.1; -;
 DR EMBL; AF527418; AM77697.1; -;
 DR PIR; A36048; A41275;
 DR Genew; HGNC:6598; LIG1.
 DR MIM; 126391; -;
 DR GO; GO:0005634; C:nucleus; TAS.
 DR GO; GO:0003677; F:DNA binding; TAS.
 DR GO; GO:0003909; F:DNA ligase activity; TAS.
 DR GO; GO:0006281; P:DNA repair; TAS.
 DR GO; GO:0007345; P:embryogenesis and morphogenesis; TAS.
 DR InterPro; IPR000977; DNA_ligase.
 DR Pfam; PF01068; DNA_ligase_1.
 DR Pfam; PF04679; DNA_ligase_A_C; 1.
 DR Pfam; PF04675; DNA_ligase_A_W; 1.
 DR TIGRfam; TIGR00574; dnl1; 1.
 DR PROSITE; PS00697; DNA_LIGASE_A1; 1.
 DR PROSITE; PS00333; DNA_LIGASE_A2; 1.
 DR PROSITE; PS0160; DNA_LIGASE_A3; 1.
 KW ATP-binding; Nuclear protein; DNA replication; Cell division; Ligase;
 KW Disease mutation; Phosphorylation; Polymorphism;
 FT BINDING 568 AMP (BY SIMILARITY).
 FT VARIANT 249 G -> E (in dbSNP:3730911).
 FT VARIANT 267 N -> S (in dbSNP:3730933).
 FT VARIANT 409 R -> H (in dbSNP:4987068).
 FT VARIANT 480 M -> V (in dbSNP:3730980).
 FT VARIANT 566 E -> K (in LIG1 deficiency).
 FT VARIANT 614 T -> I (in dbSNP:3731003).
 FT VARIANT 771 R -> W (in LIG1 deficiency).
 FT VARIANT 771 /FTID=VAR_002263.
 SQ SEQUENCE 919 AA; 101735 MW; B2854D3E38A8D4D CRC64;
 Query Match 100.0%; Score 29; DB 1; Length 919;
 Best Local Similarity 100.0%; Pred. No. 2.1e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPR 5

Db 218 TKPR 222

RESULT 13

SECS_DROME
 ID SECS_DROME STANDARD; PRT; 985 AA.
 AC Q9VNH6; 2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 42, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Probable exocyst complex component Sec8.
 GN CG2095.
 OS Drosophila melanogaster (fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Berkley;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celinker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.B., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.C., Wortman J.R., Yandell M.D., Zhang Q., Chen L.Y.,
 RA Brandon R.C., Rogers Y.-C., Blazer B.G., Heit G., Nelson C.R., Miklos G.L.G.,
 RA Wan K.H., Doyle C., Baxter B.G., Helt G., Andrews-Pfankoch C., Baldwin D.,
 RA Abrell J.F., Aghayani A., An H.-J., Andrews-Pfankoch C., Baldwin D.,
 RA Bailew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley S.M.,
 RA Beeson K.Y., Beron P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brottier P.,
 RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Recha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
 RA Foster C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D.A., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwan C.,
 RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lakso P., Lei Y., Levitsky A.A., Li J.H., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy E., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
 RA Palazolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wassarman D.A., Weinstein G.M., Weissbach J., Yao Q.A.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu S., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster";
 RL Science 287:2185-2195(2000).
 RN [2]
 RP REVISIONS.
 RX MEDLINE=22426069; PubMed=12537572;
 RA Misra S., Crosby M.A., Mungall C.J., Matthews B.B., Campbell K.S.,
 RA Hradecky P., Huang Y., Kaminker J.S., Millburn G.H., Prochnik S.E.,
 RA Smith C.D., Tupy J.L., Whitfield E.J., Bayraktaroglu L., Berman B.P.,
 RA Bettencourt B.R., Celinker S.E., de Grey A.D.N.J., Drysdale R.A.,
 RA Harris N.L., Richter J., Russo S., Schroeder A.J., Shu S.Q.,
 RA Stapleton M., Yamada C., Ashburner M., Gelbart W.M., Rubin G.M.,
 RA Lewis S.E.;
 RT "Annotation of the Drosophila melanogaster euchromatic genome: a
 RT systematic review";
 RL Genome Biol. 3:RESEARCH0083.1-RESEARCH0083.22(2002).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Berkley; TISSUE=Embryo;
 RX MEDLINE=22426066; PubMed=12537569;
 RA Stapleton M., Carlson J.W., Brokstein P., Yu C., Champe M.,

George R.A., Guarin H., Kronmiller B., Pacleb J.M., Park S., Wan K.H.,
 RA Rubin G.M., Celniker S.E.,
 RA "A Drosophila full-length cDNA resource";
 RL Genome Biol. 3:RESEARCH0080.1-RESEARCH0080.8(2002).
 CC -!- FUNCTION: Component of the exocyst complex involved in the docking
 CC of exocyst vesicles with fusions site on the plasma membrane (By
 CC similarity).
 CC SUBUNIT: The exocyst complex is composed of SEC3, SEC5, SEC6,
 CC SEC8, SEC10, SEC15, EXO70 and EXO84 (By similarity).
 CC -!- SIMILARITY: Belongs to the SEC8 family.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (see <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: A5003601; AAF51959.3; --
 CC EMBL: AY119660; AAM50314.1; --
 CC FlyBase: FBgn0037373; CG2095.
 CC InterPro: IPR004172; L27.
 CC InterPro: IPR007191; Sec8_exocyst.
 CC Pfam: PF04048; Sec8_exocyst; 1.
 CC Pfam: PF02828; L27; 1.
 CC DR Exocytosis; Transport; Protein transport; Coiled coil.
 CC KW EXOCYTOSIS; TRANSPORT; COILED COIL (POTENTIAL).
 CC FT DOMAIN 36 70
 CC CONFLICT 118 118 M -> T (IN REF. 3).
 CC CONFLICT 209 209 E -> D (IN REF. 3).
 CC CONFLICT 333 333 S -> T (IN REF. 3).
 CC CONFLICT 668 668 Y -> S (IN REF. 3).
 CC FT CONFLICT 668 668 Y -> S (IN REF. 3).
 CC FT CONFLICT 985 AA; 111665 MW; 6FBP0D9C539FBE7 CRC64;
 CC SQ
 Query Match 100.0%; Score 29; DB 1; Length 985;
 Best Local Similarity 100.0%; Pred. No. 2.3e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 7 TKPPR 11
 RESULT 14
 ID BAB2 DROME STANDARD; PRT; 1067 AA.
 AC Q9W0K4; Q24001; Q9UH3;
 DT 10-OCT-2003 (Rel. 42, Created)
 DT 10-OCT-2003 (Rel. 42, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Bric-a-brac protein 2.
 GN BAB2 OR BVBI OR CG9102/CG13911.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Hexapoda; Insecta; Pterygota;
 OC Ephydroidea; Drosophilidae; Diptera; Brachycera; Muscomorpha;
 OC NCR1_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A., FUNCTION, SUBCELLULAR LOCATION, AND TISSUE
 RP SPECIFICITY.
 RC TISSUE=Embryo, and Ovary;
 RX MEDLINE=21969340; PubMed=11973274;
 RA Couderc J.L.G., Godt D., Zolman S., Chen J., Li M., Tjong S.,
 RA Cranton S.E., Sanut-Barnola I., Laski F.A.;
 RT "The bric-a-brac locus consists of two paralogous genes encoding
 RT BTB/POZ domain proteins and acts as a homeotic and morphogenetic
 RT regulator of imaginal development in Drosophila";
 RL Development 129:2419-2433(2002).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Berkeley;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,

Ananides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.C., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon K.C., Rogers Y.-H.C., Blazer R.G., Champagne M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkuch C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brottier P.,
 RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.B., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
 RA Fesler K., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
 RA Glodok A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D.A., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
 RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lasko P., Lei Y., Levitsky A.A., Li J.H., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Swirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster";
 RL Science 287:2185-2195(2000).
 RN [3]
 RP REVISIONS.
 RX MEDLINE=22426069; PubMed=12537572;
 RA Misra S., Crosby M.A., Mungall C.J., Matthews B.B., Campbell K.S.,
 RA Hradecky P., Huang Y., Kaminker J.S., Millburn G.H., Prochuk S.E.,
 RA Smith C.D., Tupy J.L., Whitfield E.J., Bayraktaroglu L., Berman B.P.,
 RA Bettencourt B.R., Celniker S.E., de Grey A.D.N.J., Drysdale R.A.,
 RA Harris N.L., Richter J., Russo S., Schroeder A.J., Shu S.Q.,
 RA Stapleton M., Yamada C., Ashburner M., Gelbart W.M., Rubin G.M.,
 RA Lewis S.E.;
 RT "Annotation of the Drosophila melanogaster euchromatic genome: a
 RT systematic review";
 RL Genome Biol. 3:RESEARCH0083.1-RESEARCH0083.22(2002).
 RN [4]
 RP SEQUENCE OF 196-310 FROM N.A.
 RX MEDLINE=95024186; PubMed=7938017;
 RA Zolman S., Godt D., Prive G.G., Couderc J.L., Laski F.A.;
 RT "The BTB domain, found primarily in zinc finger proteins, defines an
 RT evolutionarily conserved family that includes several developmentally
 RT regulated genes in Drosophila";
 RL Proc. Natl. Acad. Sci. U.S.A. 91:10717-10721(1994).
 CC -!- FUNCTION: Probably acts as a transcriptional regulator. Required
 CC for the specification of the tarsal segment. Also involved in
 CC antenna development.
 CC -!- SUBCELLULAR LOCATION: Nuclear.
 CC -!- TISSUE SPECIFICITY: Leg imaginal disk at the central region of the
 CC tarsus and in eye antenna disk at the basal cylinder.
 CC -!- MISCELLANEOUS: 'Bric-a-brac' means 'jumble' in French (referring to
 CC the mutant ovary phenotype).
 CC -!- SIMILARITY: Contains 1 A.T hook DNA-binding repeat.
 CC -!- SIMILARITY: Contains 1 BTB/POZ domain.
 CC -!- SIMILARITY: Contains 1 helix-turn-helix Bq-type domain.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its

CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).

DR EMBL; AJ252173; CAB64388.1; -;
 DR EMBL; AEO03470; AAF47442.2; -;
 DR EMBL; U14399; AAS0834.1; -;
 DR Flybase; F590025525; bab2.
 DR InterPro; IPR000637; AT hook.
 DR InterPro; IPR000210; BTB_POZ.
 DR Pfam; PF00651; BTB; 1.
 DR SMART; SM00225; BTB; 1.
 DR PROSITE; PS00097; BTB; 1.
 DR Nuclear protein; DNA-binding; Transcription regulation.
 FT DOMAIN 223 288
 FT DNA_BIND 645 690 H-T-H MOTIF PSQ-TYPE.
 FT DNA_BIND 697 708 A-T HOOK.
 FT CONFLICT 858 858 A -> R (IN REF. 1).
 SQ SEQUENCE 1067 AA; 114661 MW; 7DBFC7681D507FC0 CRC64;

Query Match 100.0%; Score 29; DB 1; Length 1067;
 Best Local Similarity 100.0%; Pred. No. 2.5e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TKPDR 5
 Db 181 TKPDR 185

RESULT 15
 ID -TRK1 YEAST STANDARD; PRT; 1235 AA.
 AC P12685;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 01-OCT-1989 (Rel. 12, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Potassium transport protein, high-affinity.
 GN TRK1 OR YJL129C OR J0693.
 OS Saccharomyces cerevisiae (Baker's Yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
 OX NCBI_TaxID=4932;
 [1]
 SEQUENCE FROM N.A.
 RX MEDLINE=88302204; PubMed=3043197;
 RA Gaber R.F., Styles C.A., Fink G.R.;
 RT "TRK1 encodes a plasma membrane protein required for high-affinity
 RT potassium transport in *Saccharomyces cerevisiae*.";
 EL Mol. Cell. Biol. 8:2848-2859(1988).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C / FY1679;
 RX MEDLINE=97103775; PubMed=8948101;
 RA Cziepluch C., Kordes E., Pujoil A., Jauniaux J.-C.;
 RT "Sequencing analysis of a 40.2 kb fragment of yeast chromosome X
 RT reveals 19 open reading frames including URA2 (5' end), TRK1, PBS2,
 RT SP10, GDI1, RPE1, PHO86, NCA3, ASE1, COT7, GZF3, two tRNA genes,
 RT three remnant delta elements and a Ty4 transposon.";
 RL Yeast 12:1471-1474(1996).
 RC -1- FUNCTION: This protein is required for high-affinity potassium
 RC transport.

CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- SIMILARITY: Belongs to the trkH potassium transport family.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).

DR EMBL; M21328; AAA34728.1; -;
 DR EMBL; Z49404; CAA89424.1; -;
 DR PIR; S05849; PWBTH.
 DR Germonline; 141741; -;
 DR SGD; S0003665; TRK1.
 DR GO; GO:0015079; P:potassium ion transporter activity; IDA.
 DR GO; GO:0030007; P:potassium ion homeostasis; IDA.
 DR InterPro; IPR003445; Cat_transp.
 DR InterPro; IPR004773; Ktransp_euk.
 DR Pfam; PF02386; TrkH; 1.
 DR TIGRFAMs; TIGR002450; K+ transporter TRK; 1.
 DR TIGRFAMs; TIGR00934; 2a38euk; 1.
 FT TRANSMEM 49 70 POTENTIAL.
 FT TRANSMEM 78 98 POTENTIAL.
 FT TRANSMEM 107 127 POTENTIAL.
 FT TRANSMEM 778 800 POTENTIAL.
 FT TRANSMEM 813 834 POTENTIAL.
 FT TRANSMEM 838 858 POTENTIAL.
 FT TRANSMEM 862 882 POTENTIAL.
 FT TRANSMEM 898 918 POTENTIAL.
 FT TRANSMEM 923 943 POTENTIAL.
 FT TRANSMEM 971 991 POTENTIAL.
 FT TRANSMEM 1078 1098 POTENTIAL.
 FT TRANSMEM 1111 1131 POTENTIAL.
 FT CARBOHYD 100 100 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 169 169 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 222 222 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 227 227 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 251 251 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 369 369 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 383 383 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 497 497 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 501 501 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 532 532 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 580 580 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 677 677 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 919 919 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 1030 1030 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 1135 1135 N-LINKED (GLCNAC. .) (POTENTIAL).
 SQ SEQUENCE 1235 AA; 141072 MW; BCE9FD8A0BA0982B CRC64;

Query Match 100.0%; Score 29; DB 1; Length 1235;
 Best Local Similarity 100.0%; Pred. No. 2.9e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TKPDR 5
 Db 438 TKPDR 442

RESULT 16
 ID -TRK1 SACBA STANDARD; PRT; 1241 AA.
 AC P28569;
 DT 01-DEC-1992 (Rel. 24, Created)
 DT 01-DEC-1992 (Rel. 24, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Potassium transport protein, high-affinity.
 GN TRK1.
 OS Saccharomyces bayanus (Yeast) (Saccharomycetes uvarum).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
 OX NCBI_TaxID=4931;
 [1]
 SEQUENCE FROM N.A.
 RC STRAIN=R1668;
 RX MEDLINE=91216443; PubMed=2023232;
 RA Anderson J.A., Best L.A., Gaber R.F.;
 RT "Structural and functional conservation between the high-affinity K+
 RT transporters of *Saccharomyces uvarum* and *Saccharomyces cerevisiae*.";
 RL Gene 99:39-46(1991).
 CC -1- FUNCTION: This protein is required for high-affinity potassium

transport.
 CC -!- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -!- SIMILARITY: Belongs to the trkH potassium transport family.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; M57508; AAA34661.1; -;
 CC InterPro; IPR003445; Cat. transp.
 CC InterPro; IPR004773; Ktransp_euk.
 CC Pfam; PF02386; TrkH; 1.
 CC PIRSF; PIRSF002450; K. transp. TRK; 1.
 CC TIGRFAMs; TIGR00934; 2a38euk; 1.
 CC KW TRANSPORT; Transmembrane; Potassium transp.; Glycoprotein.
 CC FT TRANSMEM 49 70 POTENTIAL.
 CC FT TRANSMEM 78 98 POTENTIAL.
 CC FT TRANSMEM 107 127 POTENTIAL.
 CC FT TRANSMEM 784 806 POTENTIAL.
 CC FT TRANSMEM 819 840 POTENTIAL.
 CC FT TRANSMEM 844 864 POTENTIAL.
 CC FT TRANSMEM 868 888 POTENTIAL.
 CC FT TRANSMEM 904 924 POTENTIAL.
 CC FT TRANSMEM 929 949 POTENTIAL.
 CC FT TRANSMEM 977 997 POTENTIAL.
 CC FT TRANSMEM 1084 1104 POTENTIAL.
 CC FT TRANSMEM 1117 1137 POTENTIAL.
 CC FT CARBOHYD 100 100 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 223 223 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 227 227 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 233 233 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 257 257 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 274 274 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 353 353 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 364 364 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 389 389 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 442 442 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 505 505 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 538 538 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 584 584 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 660 660 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 681 681 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 691 691 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 741 741 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 925 925 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC FT CARBOHYD 1141 1141 N-LINKED (GLCNAC. .) (POTENTIAL).
 CC SEQUENCE 1241 AA; 141225 MW; F5149E47EC00BCD2 CRC64;
 Query Match 100.08; Score 29; DB 1; Length 1241;
 Best Local Similarity 100.08; Pred. No. 2.9e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 Db 444 TKPPR 448
 RESULT 17
 RS14 PYRAE STANDARD; PRT; 54 AA.
 AC Q8ZVM1; P58732;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE 30S ribosomal protein S14p.
 GN RPS14P OR PAE2097.
 OS Pyrobaculum aerophilum.
 OC Archaea; Crenarchaeota; Thermoprotei; Thermoproteales;
 OC Thermoproteaceae; Pyrobaculum.

NCBI_TaxID=13773;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=IM2 / ATCC 51768 / DSM 7523;
 RX MEDLINE=21664397; PubMed=11792869;
 RA Fitz-Gibbon S.R.; Ladner H., Kim U.-J., Stetter K.O., Simon M.I.,
 RA Miller J.H.;
 RT "Genome sequence of the hyperthermophilic crenarchaeon Pyrobaculum
 aerophilum";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:984-989(2002).
 CC -!- SIMILARITY: Belongs to the S14p family of ribosomal proteins.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; AE009857; AAL63943.1; -;
 CC InterPro; IPR001209; Ribosomal_S14.
 CC Pfam; PF00253; Ribosomal_S14; 1.
 CC PROSITE; PS00527; RIBOSOMAL_S14; 1.
 CC Ribosomal protein; Complete Proteome.
 CC KW RIBOSOMAL PROTEIN; Complete Proteome.
 CC SEQUENCE 54 AA; 6444 MW; 153B7BB801EDD963 CRC64;
 Query Match 89.78; Score 26; DB 1; Length 54;
 Best Local Similarity 80.08; Pred. No. 47;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 Db 4 TKPPK 8
 RESULT 18
 SECR MOUSE STANDARD; PRT; 133 AA.
 ID Q09535;
 AC Q09535; (Rel. 30, Created)
 DT 01-OCT-1994 (Rel. 30, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Secretin precursor.
 GN SCT.
 OS Mus musculus (Mouse).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 CC NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=94234995; PubMed=8179583;
 RA Lan M.S., Kajiyama W., Donadel G., Lu J., Notkins A.L.;
 RT "cDNA sequence and genomic organization of mouse secretin";
 RL Biochem. Biophys. Res. Commun. 200:1066-1071(1994).
 CC -!- FUNCTION: Stimulates formation of NaHCO(3)-rich pancreatic juice
 CC and secretion of NaHCO(3)-rich bile and inhibits HCl production by
 CC the stomach.
 CC -!- SUBCELLULAR LOCATION: Secreted.
 CC -!- SIMILARITY: Belongs to the glucagon family.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; U07568; AAA18453.1; -;
 CC EMBL; X73580; CAA51982.1; -;
 CC PIR; JC2202; JC2202.
 CC MGD; MGI:99466; Sct.

DR InterPro; IPR000532; Glucagon.
 DR Pfam; PF00123; hormone2; 1.
 DR SMART; SM00070; GLUCA; 1.
 DR PROSITE; PS00260; GLUCAGON; 1.
 KW Glucagon family; Hormone; Amidation;
 FT SIGNAL 1 22 BY SIMILARITY.
 FT PEPTIDE 32 58 SECRETIN (BY SIMILARITY).
 FT MOD_RES 58 58 AMIDATION (G-59 PROVIDE AMIDE GROUP).
 SQ SEQUENCE 133 AA; 14914 MW; 9869CBF74CA9709 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 133;
 Best Local Similarity 80.0%; Pred. No. 1.2e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 DB 129 TRPRP 133

RESULT 19

VEGH ORFN2 STANDARD; PRT; 133 AA.

AC P525B4; (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Vascular endothelial growth factor homolog precursor.
 GN A28.

OS Orf virus (strain NZ2) (OV NZ-2).
 OC Viruses; dsDNA viruses, no RNA stage; Poxviridae; Chordopoxvirinae;
 OC Parapoxvirus.
 OC NCBI_TaxID=10259;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=94076465; PubMed=8254780;

RA Lytle D.J., Fraser K.M., Fleming S.B., Mercer A.A., Robinson A.J.;
 RT "Homologs of vascular endothelial growth factor are encoded by the
 RT poxvirus orf virus."
 RL J. Virol. 68:84-92(1994).

CC -!- FUNCTION: INDUCES ENDOTHELIAL PROLIFERATION.

CC -!- SUBUNIT: Homodimer; disulfide-linked (BY SIMILARITY).

CC -!- SIMILARITY: Belongs to the PDGF/VEGF growth factor family.

CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).

CC -----

DR EMBL; S67520; AAB29220.2; --

DR HSSP; P15692; 1VPP.

DR InterPro; IPR002400; GF_cysknot.

DR InterPro; IPR000072; PD_growth_factor.

DR Pfam; PF00341; PDGF; 1.

DR PRINTS; PR00438; GFCSKNOT.

DR PRODOM; PD001629; PD_growth_factor; 1.

DR SMART; SM00141; PDGF; 1.

DR PROSITE; PS00249; PDGF_1; 1.

DR PROSITE; PS02078; PDGF_2; 1.

KW Mitogen; Growth factor; Glycoprotein; Signal.

FT SIGNAL 1 20 POTENTIAL.

FT CHAIN 21 133 VASCULAR ENDOTHELIAL GROWTH FACTOR

FT HOMOLOG.

FT DISULFID 36 78 BY SIMILARITY.

FT DISULFID 67 112 BY SIMILARITY.

FT DISULFID 71 114 BY SIMILARITY.

FT DISULFID 61 61 INTERCHAIN (BY SIMILARITY).

FT DISULFID 70 70 INTERCHAIN (BY SIMILARITY).

FT CARBOHYD 85 85 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT SEQUENCE 133 AA; 14715 MW; 917C0F6883030C39 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 133;
 Best Local Similarity 80.0%; Pred. No. 1.2e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 DB 126 TRPRP 130

RESULT 20

RT12 DROME STANDARD; PRT; 140 AA.

AC P10735; Q9V3R2;

DT 01-JUL-1989 (Rel. 11, Created)

DT 01-JUL-1989 (Rel. 11, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)

DE 40S ribosomal protein S12, mitochondrial precursor (MT-RPS12)

DE (technical knockout locus protein).

GN TKO OR EG:BACH5J11.1 OR CG7925.

OS Drosophila melanogaster (Fruit fly).

OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;

OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;

OC Ephydroidea; Drosophilidae; Drosophila.

OC NCBI_TaxID=7227;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=88027001; PubMed=3117373;

RA Royden C.S., Pirrotta V., Jan L.Y.;

RT "The tko locus, site of a behavioral mutation in D. melanogaster,
 RT codes for a protein homologous to prokaryotic ribosomal protein
 RT S12."
 RL Cell 51:165-173(1987).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=Oregon-R;

RX MEDLINE=20196011; PubMed=10731137;

RA Benos P.V., Gatt M.K., Ashburner M., Murphy L., Harris D.,

RA Barrell B.G., Ferraz C., Vidal S., Brun C., Demailles J., Cadieu B.,

RA Dreano S., Gloux S., Lelaure V., Mottier S., Galibert F., Borkova D.,

RA Papagannakis G., Spanos L., Cox S., Madueno E., de Pablo B.,

RA Modoll J., Peter A., Schoettler P., Werner M., Moutkioti F.,

RA Benister D.M., Campbell L.A., Darlamitsova A., Henderson N.S.,

RA McWilliam P.J., Salles C., Tait E.A., Valenti P., Saunders R.D.C.,

RA Glover D.M.;

RT "From sequence to chromosome: the tip of the X chromosome of D.
 RT melanogaster."
 RL Science 287:2220-2222(2000).

RN [3]

RP SEQUENCE FROM N.A.

RC STRAIN=Berkley;

RX MEDLINE=20196006; PubMed=10731132;

RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,

RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,

RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,

RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,

RA Brandon R.C., Rogers Y.-H.C., Blaise R.G., Champ M., Pfeiffer B.D.,

RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,

RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfankuch C., Baldwin D.,

RA Bailew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,

RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,

RA Borkova D., Botchan M.R., Botkin J., Brokstein P., Brottier P.,

RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,

RA de Pablo B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,

RA Dodson K.J., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,

RA Fozler C., Gabriellian A.E., Garg N.S., Gelbart W.M., Glasser K.,

RA Glodex A., Gong F., Gorrell J.H., Gu Z., Guan Z., Harris M.,

RA Harris N.I., Harvey D.A., Heiman T.J., Hernandez J.R., Houck J.,

RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibsen G.,

```
RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Laslo P., Lei Y., Levitsky A.A., Li J.H., Li Z., Liang Y., Lin X.,
RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Mekullov G., Milshina N.V., Mobarri C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M.I., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinet K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Rhue B.C., Siden-Klamis I., Simpson M., Skupski M.P., Smith T.,
RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RT Science 287:2185-2195(2000).
CC -!- SUBCELLULAR LOCATION: Mitochondrial.
CC -!- MISCELLANEOUS: MUTATION OF TKO CAUSES BEHAVIORAL MUTATION ("BANG
CC SENSITIVITY" = TEMPORARILY PARALYSIS IN RESPONSE TO A PHYSICAL
CC JOLT).
CC -!- SIMILARITY: Belongs to the S12P family of ribosomal proteins.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; M19494; AAA28935.1; -.
CC EMBL; AL33505; CAB65841.1; -.
CC EMBL; AE003424; AAF45781.1; -.
CC PIR; A29622; A29622.
CC DR FlyBase; FBgn0003714; tko.
CC DR GO; GO:0008049; P: male courtship behavior; IMP.
CC DR GO; GO:0007638; P: mechanosensory behavior; IMP.
CC DR GO; GO:0009592; P: perception of sound; IMP.
CC DR GO; GO:00095612; P: response to mechanical stimulus; IMP.
CC DR InterPro; IPR008994; Nucleic acid OB.
CC DR InterPro; IPR006032; Ribosomal S12.23.
CC DR InterPro; IPR005679; Ribosomal S12b/c.
CC DR Pfam; PF00164; Ribosomal S12; I.
CC DR PRINTS; PF01034; RIBOSOMALS12.
CC DR PRODOM; PD000576; Ribosomal_S12_23; 1.
CC DR TIGRFAMs; TIGR00981; rpsL_bact; 1.
CC DR PROSITE; PS00055; RIBOSOMAL_S12; 1.
CC KW Ribosomal protein; Mitochondrion; Transit peptide.
CC FT TRANSIT 1 30 MITOCHONDRION (POTENTIAL).
CC FT CHAIN 31 140 40S RIBOSOMAL PROTEIN S12.
CC SQ SEQUENCE 140 AA; 15532 MW; 8187A7312C529F31 CRC64;
Query Match 89.7%; Score 26; DB 1; Length 140;
Best Local Similarity 80.0%; Pred. No. 1.2e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPRP 5
DB 45 TRPRP 49
RESULT 21
KGUA SYNEL
ID KGUA SYNEL STANDARD; PRT; 191 AA.
AC Q8DMQ7;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Guanylate kinase (EC 2.7.4.8) (GMP kinase).
GN GMK OR TLL0054.
```

```
OS Synechococcus elongatus (Thermosynechococcus elongatus).
OC Bacteria; Cyanobacteria; Chroococcales; Synechococcus.
CX NCBI_TaxID=32046;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=SP-1;
RX MEDLINE=222525144; PubMed=12240834;
RA Nakamura Y., Kaneo T., Sato S., Ikeuchi M., Katoh H., Sasamoto S.,
RA Watanabe A., Iriguchi M., Kawashima K., Kimura T., Kishida Y.,
RA Kiyokawa C., Kohara M., Matsumoto M., Matsuno A., Nakazaki N.,
RA Shimpo S., Sugimoto M., Takeuchi C., Yamada M., Tabata S.;
RT "Complete genome structure of the thermophilic cyanobacterium
RT Thermosynechococcus elongatus SP-1.";
RL DNA Res. 9:123-130(2002).
CC -!- FUNCTION: Essential for recycling GMP and indirectly, cGMP.
CC -!- CATALYTIC ACTIVITY: ATP + GMP = ADP + GDP.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the guanylate kinase family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AP005369; BAC07607.1; -.
CC DR HAMAP; MF 00328; -.
CC DR InterPro; IPR008144; Guanylate_kin.
CC DR InterPro; IPR008145; Guanylt/Ca.
CC DR Pfam; PF00625; Guanylate_kin; 1.
CC DR SMART; SM00072; GuKc; 1.
CC DR PROSITE; PS00856; GUANYLATE_KINASE_1; 1.
CC DR PROSITE; PS00052; GUANYLATE_KINASE_2; 1.
CC KW Transferase; Kinase; ATP-binding; Complete proteome.
CC FT NP BIND 16 23 ATP (BY SIMILARITY).
CC SQ SEQUENCE 191 AA; 20990 MW; 70B6C15768D73D51 CRC64;
Query Match 89.7%; Score 26; DB 1; Length 191;
Best Local Similarity 80.0%; Pred. No. 1.7e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPRP 5
DB 44 TRPRP 48
RESULT 22
VCO7 ADE04
ID VCO7 ADE04 STANDARD; PRT; 193 AA.
AC Q96831;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Major core protein precursor (Protein VII) (pVII).
GN PVII.
OS Human adenovirus type 4.
OC Viruses; dsDNA viruses, no RNA stage; Adenoviridae; Mastadenovirus.
CX NCBI_TaxID=28280;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Isolate RI-6;
RA Tarassishin L., Swalowski P.W.S., McIay J., Russell W.C.;
RL Submitted (OCT-1996) to the EMBL/GenBank/DBJ databases.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
```

```
CC EMBL; U70921; AAC83411.1; -.
DR InterPro; IPR004912; Adeno_VII.
DR Pfam; PF03228; Adeno_VII; 1.
FT PROPEP 1 24 BY SIMILARITY.
FT CHAIN 25 193 MAJOR CORE PROTEIN.
FT SITE 24 25 CLEAVAGE (BY ADENOVIRUS PROTEASE)
FT SITE 24 25 (POTENTIAL).
SQ SEQUENCE 193 AA; 21358 MW; 43137E07DB379DD0 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 193;
Best Local Similarity 80.0%; Pred. No. 1.7e+02;
Matches 4; Conservative 1; Mismatches 0; Gaps 0;

Qy 1 TKPRP 5
Db 188 TRPPR 192

RESULT 23
VC07_ADE02 STANDARD; PRT; 198 AA.
AC P03266; P12542;
DT 21-JUL-1986 (Rel. 01, Created)
DT 01-FEB-1994 (Rel. 28, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Major core protein precursor (Protein VII) (pVII).
GN pVII.
OS Human adenovirus type 2, and
OS Human adenovirus type 5.
OC Viruses; dsDNA viruses, no RNA stage; Adenoviridae; Mastadenovirus.
ON NCBI_TaxID=10515, 28285;
RX MEDLINE=85054835; PubMed=6094534;
RA Alestroem P., Akusjlaervi G., Lager M., Yeh-Kai L., Pettersson U.;
RT "Genes encoding the core proteins of adenovirus type 2.";
RN J Biol. Chem. 259:13980-13985(1984).
SQ SEQUENCE FROM N.A.
RC SPECIES=Human adenovirus type 2;
RX MEDLINE=85054835; PubMed=6094534;
RA Alestroem P., Akusjlaervi G., Lager M., Yeh-Kai L., Pettersson U.;
RT "Genes encoding the core proteins of adenovirus type 2.";
RN J Biol. Chem. 259:13980-13985(1984).
SQ SEQUENCE FROM N.A.
RC SPECIES=Human adenovirus type 2;
RX MEDLINE=83221511; PubMed=6574459;
RA Sung M.T., Cao T.M., Coleman R.T., Budelier K.A.;
RT "Gene and protein sequences of adenovirus protein VII, a hybrid basic chromosomal protein.";
RN Proc. Natl. Acad. Sci. U.S.A. 80:2902-2906(1983).
SQ SEQUENCE FROM N.A.
RC SPECIES=Human adenovirus type 5;
RX MEDLINE=92087470; PubMed=1727603;
RA Chroboczek J., Bieber F., Jacrot B.;
RT "Determination of the nucleotide sequence for the penton-base gene of human adenovirus type 5.";
RN Gene 69:153-157(1988).
CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).
DR EMBL; J01917; AAA92212.1; -.
DR InterPro; IPR004912; Adeno_VII.
DR Pfam; PF03228; Adeno_VII; 1.
FT PROPEP 1 24 BY SIMILARITY.
FT CHAIN 25 193 MAJOR CORE PROTEIN.
FT SITE 24 25 CLEAVAGE (BY ADENOVIRUS PROTEASE)
FT SITE 24 25 (POTENTIAL).
SQ SEQUENCE 193 AA; 21358 MW; 43137E07DB379DD0 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 193;
Best Local Similarity 80.0%; Pred. No. 1.7e+02;
Matches 4; Conservative 1; Mismatches 0; Gaps 0;

Qy 1 TKPRP 5
Db 188 TRPPR 192
```

```
DR EMBL; M73260; AAA96408.1; ALT_SEQ.
DR EMBL; M22141; AAA42520.1; -.
DR PIR; C03837; FOAD72.
DR PIR; P0067; FOADH5.
DR InterPro; IPR004912; Adeno_VII.
DR Pfam; PF03228; Adeno_VII; 1.
FT CHAIN 25 193 MAJOR CORE PROTEIN.
FT SITE 24 25 CLEAVAGE (BY ADENOVIRUS PROTEASE)
FT SITE 24 25 (POTENTIAL).
SQ SEQUENCE 193 AA; 21992 MW; 7D5ASD426F08E952 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 198;
Best Local Similarity 80.0%; Pred. No. 1.8e+02;
Matches 4; Conservative 1; Mismatches 0; Gaps 0;

Qy 1 TKPRP 5
Db 193 TRPPR 197

RESULT 24
KGUA_ANASP STANDARD; PRT; 199 AA.
AC Q8Z017;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Guanylate kinase (EC 2.7.4.8) (GMP kinase).
GN GMP OR ALR0106.
OS Anabaena sp. (strain PCC 7120).
OC Bacteria; Cyanobacteria; Nostocales; Nostocaceae; Nostoc.
ON NCBI_TaxID=103690;
RX MEDLINE=21595285; PubMed=11759840;
RA Kaneko T., Nakamura Y., Wolk C.P., Kuritz T., Sasamoto S.,
RA Watanabe A., Iriguchi M., Ishikawa A., Kawashima K., Kimura T.,
RA Kishida Y., Kohara M., Matsumoto M., Matsuno A., Muraki A.,
RA Nakazaki N., Shimpou S., Sugimoto M., Takazawa M., Yamada M.,
RA Yasuda M., Tabata S.;
RT "Complete genomic sequence of the filamentous nitrogen-fixing cyanobacterium Anabaena sp. strain PCC 7120.";
RN DNA Res. 8:205-213(2001).
CC -1- FUNCTION: Essential for recycling GMP and indirectly, CGMP.
CC -1- CATALYTIC ACTIVITY: ATP + GMP = ADP + GDP.
CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -1- SIMILARITY: Belongs to the guanylate kinase family.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce/ or send an email to license@isb-sib.ch).
DR EMBL; AF003581; BAB77630.1; -.
DR PIR; AB1820; AB1820.
DR HAMAP; MF_00328; -.
DR InterPro; IPR008144; Guanylate_kin.
DR InterPro; IPR008145; Guanylt/Ca.
DR Pfam; PF00625; Guanylate_kin; 1.
DR SMART; SM00072; GUKC; 1.
DR PROSITE; PS00856; GUANYLATE_KINASE_1; 1.
DR PROSITE; PS00852; GUANYLATE_KINASE_2; 1.
DR TRANSFERASE; Kinase; ATP-binding; Complete proteome.
FT NF_BIND 27 34 ATP (BY SIMILARITY).
SQ SEQUENCE 199 AA; 22099 MW; 58634496D484850D CRC64;

Query Match 89.7%; Score 26; DB 1; Length 199;
Best Local Similarity 80.0%; Pred. No. 1.8e+02;
```

Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|
Db 55 TRPPR 59

RESULT 25
KGUA_NEIMA
ID _KGUA_NEIMA STANDARD; PRT; 205 AA.
AC Q3JT56;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Guanylate kinase (EC 2.7.4.8) (GMP kinase).
GN GMP OR NMA1919.
OS Neisseria meningitidis (serogroup A).
OC Bacteria; Proteobacteria; Betaproteobacteria; Neisseriales;
OC Neisseriaceae; Neisseria.
OX NCBI_TaxID=65699;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=22491 / Serogroup A / Serotype 4A;
RX MEDLINE=20222556; PubMed=10761919;
RA Parkhill J., Achtman M., James K.D., Bentley S.D., Churcher C.,
RA Klee S.R., Moralli G., Basham D., Brown D., Chillingworth T.,
RA Davies R.M., Davis P., Devlin K., Feltwell T., Hamlin N., Holroyd S.,
RA Jorgels K., Leather S., Moule S., Mungall K., Quail M.A.,
RA Rajandream M.A., Rutherford K.M., Simmonds M., Skelton J.,
RA Whitehead S., Spratt B.G., Barrrell B.G.;
RT "Complete DNA sequence of a serogroup A strain of Neisseria
RT meningitidis Z2491.";
RL Nature 404:502-506(2000).
CC -!- FUNCTION: Essential for recycling GMP and indirectly, cGMP.
CC -!- CATALYTIC ACTIVITY: ATP + GMP = ADP + GDP.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the guanylate kinase family.
CC
CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announcement/>
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AL162757; CAB85140.1; -.
CC F1R; G81819; G81819.
CC HSP; P15454; IGKY.
CC HAMAP; MF_00328; -; 1.
CC InterPro; IPR008144; Guanylate_kin.
CC InterPro; IPR008145; Guanylt/Ca.
CC Pfam; PF00625; Guanylate_kin; 1.
CC SMART; SM00072; GuKc; 1.
CC PROSITE; PS00856; GUANYLATE_KINASE_1; 1.
CC PROSITE; PS00852; GUANYLATE_KINASE_2; 1.
CC Transferase; Kinase; ATP-binding; Complete proteome.
KW NP_BIND 14 21 ATP (BY SIMILARITY).
FT NP_BIND 14 21
SQ SEQUENCE 205 AA; 22530 MW; 04FA07E5450C9007 CRC64;
Query Match 89.7%; Score 26; DB 1; Length 205;
Best Local Similarity 80.0%; Pred. No. 1.8e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|
Db 42 TRPPR 46

RESULT 26
KGUA_NEIMB
ID _KGUA_NEIMB STANDARD; PRT; 205 AA.
AC Q9JYB5;

DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Guanylate kinase (EC 2.7.4.8) (GMP kinase).
GN GMP OR NMA1661.
OS Neisseria meningitidis (serogroup B).
OC Bacteria; Proteobacteria; Betaproteobacteria; Neisseriales;
OC Neisseriaceae; Neisseria.
OX NCBI_TaxID=491;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MC58 / Serogroup B;
RX MEDLINE=20175755; PubMed=10710307;
RA Tettelin H., Saunders N.J., Heidelberg J., Jeffries A.C., Nelson K.E.,
RA Eisen J.A., Ketchum K.A., Hood D.W., Peden J.F., Dodson R.O.,
RA Nelson W.C., Gwinn M.L., DeBoy R., Peterson J.D., Hickey E.K.,
RA Haft D.H., Salzberg S.L., White O., Fleischmann R.D., Dougherty B.A.,
RA Mason T., Ciecko A., Parksey D.S., Blair E., Ciftone H., Clark E.B.,
RA Cotton M.D., Utterback T.R., Khouri H., Qin H., Vamathevan J.,
RA Gill J., Scarlato V., Maignani V., Pizzi M., Grandi G., Sun L.,
RA Smith H.O., Fraser C.M., Moxon E.R., Rappuoli R., Venter J.C.;
RT "Complete genome sequence of Neisseria meningitidis serogroup B strain
RT MC58.";
RL Science 287:1809-1815(2000).
CC -!- FUNCTION: Essential for recycling GMP and indirectly, cGMP.
CC -!- CATALYTIC ACTIVITY: ATP + GMP = ADP + GDP.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the guanylate kinase family.
CC
CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announcement/>
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AE002517; AAF42010.1; -.
CC F1R; G81055; G81055.
CC HSP; P15454; IGKY.
CC TIGR; NME1661; -.
CC HAMAP; MF_00328; -; 1.
CC InterPro; IPR008144; Guanylate_kin.
CC InterPro; IPR008145; Guanylt/Ca.
CC Pfam; PF00625; Guanylate_kin; 1.
CC SMART; SM00072; GuKc; 1.
CC PROSITE; PS00856; GUANYLATE_KINASE_1; 1.
CC PROSITE; PS00852; GUANYLATE_KINASE_2; 1.
CC Transferase; Kinase; ATP-binding; Complete proteome.
KW NP_BIND 14 21 ATP (BY SIMILARITY).
FT NP_BIND 14 21
SQ SEQUENCE 205 AA; 22500 MW; 54EE07E545189008 CRC64;
Query Match 89.7%; Score 26; DB 1; Length 205;
Best Local Similarity 80.0%; Pred. No. 1.8e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|
Db 42 TRPPR 46

RESULT 27
Y132_NPVAC
ID _Y132_NPVAC STANDARD; PRT; 219 AA.
AC P24730;
DT 01-MAR-1992 (Rel. 21, Created)
DT 01-MAR-1992 (Rel. 21, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE Hypothetical 25.1 kDa protein in pp34-EXO intergenic region (ORF 4).
OS Autographa californica nuclear polyhedrosis virus (AcMNPV).
OC Viruses; dsDNA viruses, no RNA stage; Baculoviridae;
OC Nucleopolyhedrovirus.
OX NCBI_TaxID=46015;

RN SEQUENCE FROM N.A.
 RP MEDLINE=97311863; PubMed=3041026;
 RA Oellig C., Happ B., Mueller T., Doerfler W.;
 RT "Overlapping sets of viral RNAs reflect the array of polypeptides in
 RT the cloverleaf J and N fragments (map positions 81.2 to 85.0) of the
 RT Autographa californica nuclear polyhedrosis virus genome.";
 RL J. Virol. 61:3048-3057(1987).
 CC [2]
 DR SEQUENCE FROM N.A.
 RP STRAIN=C6;
 RC MEDLINE=94303173; PubMed=8030224;
 RA Ayres M.D., Howard S.C., Kuzio J., Lopez-Ferber M., Possee R.D.;
 RT "The complete DNA sequence of Autographa californica nuclear
 RT polyhedrosis virus." (1994).
 RL Virology 202:586-605(1994).
 CC [1- SIMILARITY: TO CORRESPONDING ORF IN OPNVPV.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
 CC or send an email to license@isb-sib.ch).
 CC
 DR EMBL; M17548; AAA66806.1; -;
 DR EMBL; L22858; AAA66762.1; -;
 DR PIR; E72866; E72866.
 KW Hypothetical protein; Late protein.
 SQ SEQUENCE 219 AA; 25136 MW; 281E1625BE8F6A5E CRC64;
 CC
 Query Match 89.7%; Score 26; DB 1; Length 219;
 Best Local Similarity 80.0%; Pred. NO. 2e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 23 TKPPK 27
 CC
 RESULT 28
 PESS_LUCCU STANDARD; PRT; 220 AA.
 ID PESS_LUCCU
 AC Q95UE8; Q8MUP5;
 DT 10-OCT-2003 (Rel. 42, Created)
 DT 10-OCT-2003 (Rel. 42, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Peritrophin-55 precursor.
 OS Lucilia cuprina (Greenbottle fly) (Australian sheep blowfly).
 CC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 CC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Oestroidea;
 CC Calliphoridae; Lucilia.
 CC NCBI_TaxID=7375;
 CC [1]
 RN SEQUENCE FROM N.A.; SEQUENCE OF 20-57 AND 60-94, TISSUE SPECIFICITY,
 RP AND DEVELOPMENTAL STAGE.
 RC TISSUE=Larva;
 RX MEDLINE=2243280; PubMed=12535682;
 RA Tellam R.L., Vuocolo T., Eisemann C.H., Briscoe S., Riding G.A.,
 RA Elvin C.N., Pearson R.D.;
 RT "Identification of an immuno-protective mucin-like protein,
 RT peritrophin-55, from the peritrophic matrix of Lucilia cuprina
 RT larvae".
 RL Insect Biochem. Mol. Biol. 33:239-252(2003).
 CC [1- FUNCTION: May bind oligosaccharide structures.
 CC [1- TISSUE SPECIFICITY: Larval peritrophic membrane.
 CC [1- DEVELOPMENTAL STAGE: Expressed in all 3 larval instars but not
 CC adults or eggs.
 CC [1- PTM: Glycosylated.
 CC [1- SIMILARITY: Contains 1 chitin-binding type-2 domain.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration

CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
 CC or send an email to license@isb-sib.ch).
 CC
 DR EMBL; AY055470; AAL15463.1; -;
 DR EMBL; AF15826; AAM55223.1; -;
 DR InterPro; IPR002557; Chitin bind PerA.
 DR PROSITE; PS0940; CHIT_BIND_II; 1.
 KW Glycoprotein; Signal.
 FT SIGNAL 1 19
 FT CHAIN 20 220
 FT DOMAIN 33 95
 FT CARBOHYD 29 29
 FT CONFLICT 67 67
 FT CONFLICT 69 69
 FT CONFLICT 142 142
 FT CONFLICT 164 164
 FT CONFLICT 190 190
 FT CONFLICT 190 190
 FT SEQUENCE 220 AA; 23535 MW; 88C74ED57F2ED7C7 CRC64;
 SQ
 Query Match 89.7%; Score 26; DB 1; Length 220;
 Best Local Similarity 80.0%; Pred. NO. 2e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 155 TKPPK 159
 CC
 RESULT 29
 NOCT_RAT STANDARD; PRT; 253 AA.
 ID NOCT_RAT
 AC Q9ET55;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Nocturnin (CCR4 protein homology) (Fragment).
 GN CCRN4L OR NOC OR CCR4.
 OS Rattus norvegicus (Rat).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 CC NCBI_TaxID=10116;
 CC [1]
 RN SEQUENCE FROM N.A.
 RC TISSUE=Retina;
 RA Wang Y., Osterbur D.L., Green C.B., Besharse J.C.;
 RT "Mammalian homologs of Xenopus nocturnin: conservation of structure
 RT and circadian regulation.";
 RL Submitted (OCT-1999) to the EMBL/GenBank/DBJ databases.
 CC [1- FUNCTION: Component of the circadian clock or downstream effector
 CC of clock function. Exhibits a high amplitude circadian rhythm with
 CC maximal levels in early evening. In constant darkness or constant
 CC light, the amplitude of the rhythm decreases (By similarity).
 CC [1- SIMILARITY: Belongs to the CCR4/nocturnin family.
 CC
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
 CC or send an email to license@isb-sib.ch).
 CC
 DR EMBL; AF199495; AAG01390.1; -;
 DR InterPro; IPR005135; Exo_endo_phos.
 DR Pfam; PF03372; Exo_endo_phos; 1.
 KW Biological rhythms.
 FT NON_TER 1 1
 FT NON_TER 253 253
 FT SEQUENCE 253 AA; 28662 MW; C63566735BCC432E CRC64;
 SQ


```
Query Match      89.7%; Score 26; DB 1; Length 253;
Best Local Similarity 80.0%; Pred. No. 2.3e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
   |||
Db 44 TRPPR 48

RESULT 30
TRUA_XYLFA STANDARD; PRT; 257 AA.
AC Q9PDK6;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DE tRNA pseudouridine synthase A (EC 4.2.1.70) (Pseudouridylylate synthase
DE I) (Pseudouridine synthase I) (Uracil hydrolyase).
GN TRUA OR XF1373.
OS Xylella fastidiosa.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xylella.
OX NCBI_TaxID=2371;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=945C;
RX Simpson A.J.G., Reinach F.C., Arruda P., Abreu F.A., Acencio M.,
RA Alvares R., Alves L.M.C., Araya J.E., Baia G.S., Baptista C.S.,
RA Barros M.H., Bonaccorsi E.D., Bordin S., Bove J.M., Briones M.R.S.,
RA Bueno M.R.P., Camargo A.A., Camargo L.B.A., Carraro D.M., Carrer H.,
RA Colauto N.B., Colombo C., Costa F.F., Costa M.C.R., Costa-Neto C.M.,
RA Coutinho L.L., Cristofani M., Dias-Neto E., Docena C., El-Dorry H.,
RA Facinani A.P., Ferreira A.J.S., Ferreira V.C.A., Ferro J.A.,
RA Fraga J.S., Franca S.C., Franco M.C., Frohme M., Furlan L.R.,
RA Garnier M., Goldman G.H., Goldman M.H.S., Gomes S.L., Gruber A.,
RA Ho P.L., Hhseisel J.D., Junqueira M.L., Kemper E.L., Kitajima J.P.,
RA Krieger J.B., Kuramae E.E., Laigret F., Lambais M.R., Leite L.C.C.,
RA Lemos E.G.M., Lemos M.V.F., Lopes S.A., Lopes C.R., Machado J.A.,
RA Machado M.A., Madeira A.M.B.N., Madeira H.M.F., Marino C.L.,
RA Marques M.V., Martins E.A.L., Martins E.M.F., Matsukuma A.Y.,
RA Menck C.F.M., Miracca E.C., Miyaki C.Y., Monteiro-Vitorello C.B.,
RA Moon D.H., Nagai M.A., Nascimento A.L.T.O., Netto L.E.S.,
RA Nhani A.Jr., Nobrega F.G., Nunes L.R., Oliveira M.A.,
RA de Oliveira M.C., de Oliveira R.C., Palmeri D.A., Paris A.,
RA Peixoto B.R., Pereira G.A.G., Pereira H.A. Jr., Pequeiro J.B.,
RA Quaggio R.B., Roberto P.G., Rodrigues V., de Rosa A.J.M.,
RA de Rosa V.E. Jr., de Sa R.G., Santelli R.V., Sawasaki H.E.,
RA da Silva A.C.R., da Silva A.M., da Silva F.R., Silva W.A. Jr.,
RA da Silveira J.F., Silvestri M.L.Z., Siqueira W.J., de Souza A.A.,
RA de Souza A.P., Terenzi M.F., Truffi D., Tsai S.M., Tshako M.H.,
RA Vallada H., Van Sluys M.A., Verjovski-Almeida S., Vettore A.L.,
RA Zago M.A., Zatz M., Meidanis J., Setubal J.C.;
RT "The genome sequence of the plant pathogen Xylella fastidiosa."
RL Nature 406:151-159(2000).
CC -!- FUNCTION: Formation of pseudouridine at positions 38, 39 and 40 in
CC the anticodon stem and loop of transfer RNAs (By similarity).
CC -!- CATALYTIC ACTIVITY: Uracil + D-ribose 5-phosphate = pseudouridine
CC 5'-phosphate + H(2)O.
CC -!- SIMILARITY: Belongs to the pseudouridine synthase trUA family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AE003968; AAF64182.1; -.
CC PIR; A82691; A82691.
CC HMAP; MF_00171; -.
CC
```

```
DR InterPro: IPR001406; Pseudou synth_1.
DR Pfam; PF01416; Pseudou synth_1; 2.
DR TIGRFAMs; TIGR00071; hisf_truA; 1.
KW Lyase; tRNA processing; Complete proteome.
FT ACT_SITE 53 BY SIMILARITY.
SQ SEQUENCE 257 AA; 28638 MW; C717E0D287CD3F3 CRC64;

Query Match      89.7%; Score 26; DB 1; Length 257;
Best Local Similarity 80.0%; Pred. No. 2.3e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
   |||
Db 70 TRPPR 74

RESULT 31
TRUA_XYLFT STANDARD; PRT; 257 AA.
AC Q87DS1;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DE tRNA pseudouridine synthase A (EC 4.2.1.70) (Pseudouridylylate synthase
DE I) (Pseudouridine synthase I) (Uracil hydrolyase).
GN TRUA OR PD0610.
OS Xylella fastidiosa (strain Temecula / ATCC 700964).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xylella.
OX NCBI_TaxID=183190;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE=22421331; PubMed=12533478;
RX Van Sluys M.A., de Oliveira M.C., Monteiro-Vitorello C.B., Moon D.H.,
RA Miyaki C.Y., Furlan L.R., Camargo L.E.A., da Silva A.C.R., da Silva F.R.,
RA Takita M.A., Lemos E.G.M., Machado M.A., Ferro M.I.T., da Silva F.R.,
RA Goldman M.H.S., Goldman G.H., Lemos M.V.F., El-Dorry H., Tsai S.M.,
RA Carrer H., Carraro D.M., de Oliveira R.C., Nunes L.R., Siqueira W.J.,
RA Coutinho L.L., Kimura E.T., Ferro E.S., Harakava R., Kuramae E.B.,
RA Marino C.S., Gaglioti E., Abreu I.L., Alves L.M.C., do Amaral A.M.,
RA Baia G.S., Blanco S.R., Brito W.S., Camavari F.S., Celestino A.V.,
RA da Cunha A.F., Fenille R.C., Ferro J.A., Formighieri E.F., Kishi L.T.,
RA Leoni S.G., Oliveira A.R., Rosa V.E. Jr., Sasaki F.T., Sena J.A.D.,
RA de Souza A.A., Truffi D., Tsukumo F., Yanai G.M., Zaros L.G.,
RA Civerolo E.L., Simpson A.J.G., Almeida N.F. Jr., Setubal J.C.,
RA Kitajima J.P.;
RC "Comparative analyses of the complete genome sequences of Pierce's
RT disease and citrus variegated chlorosis strains of Xylella
RT fastidiosa."
RL J. Bacteriol. 185:1018-1026(2003).
CC -!- FUNCTION: Formation of pseudouridine at positions 38, 39 and 40 in
CC the anticodon stem and loop of transfer RNAs (By similarity).
CC -!- CATALYTIC ACTIVITY: Uracil + D-ribose 5-phosphate = pseudouridine
CC 5'-phosphate + H(2)O.
CC -!- SIMILARITY: Belongs to the pseudouridine synthase trUA family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AE012555; AAC28482.1; ALT_INIT.
CC HMAP; MF_00171; -.
CC InterPro: IPR001406; Pseudou synth_1.
DR Pfam; PF01416; Pseudou synth_1; 2.
DR TIGRFAMs; TIGR00071; hisf_truA; 1.
KW Lyase; tRNA processing; Complete proteome.
FT ACT_SITE 53 BY SIMILARITY.
SQ SEQUENCE 257 AA; 28664 MW; F6D5A76D9C2A88EC CRC64;
```

```
Query Match      89.7%; Score 26; DB 1; Length 257;
Best Local Similarity 80.0%; Pred. No. 2.3e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 70 TRPPR 74

RESULT 32
YOOB_CABEL
ID YOOB_CABEL STANDARD; PRT; 304 AA.
AC Q09360;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE Hypothetical 33.0 kDa protein EED8.11 in chromosome II precursor.
GN EED8.11.
OS Caenorhabditis elegans.
CC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
CC Rhabditidae; Peloderinae; Caenorhabditis.
CC NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Bristol N2;
RA Chissee S.;
RC SEQUENCE 304 AA; 32962 MW; 60C223B88F534151 CRC64;
RL Submitted (JUL-1995) to the EMBL/GenBank/DBSJ databases.
CC -1- SIMILARITY: SOME, TO C.ELEGANS R13F6.2 AND R13F6.8.
-----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
-----
DR EMBL; U23484; AAC46771.1; -
DR PIR; T15922; T15922.
DR WormPep; EED8.11; C01884.
DR InterPro; IPR001304; Lectin_C.
DR SMART; SM00034; CLEC7.1.
KW Hypothetical protein; Signal.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 304 HYPOTHETICAL PROTEIN EED8.11.
FT DOMAIN 64 92 POLY-THR.
SQ SEQUENCE 304 AA; 32962 MW; 60C223B88F534151 CRC64;

Query Match      89.7%; Score 26; DB 1; Length 304;
Best Local Similarity 80.0%; Pred. No. 2.7e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 51 TKPPK 55

RESULT 33
YB04_AQUAE
ID YB04_AQUAE STANDARD; PRT; 350 AA.
AC Q87189;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hypothetical protein AQ_1104.
GN AQ_1104.
OS Aquifex aeolicus.
CC Bacteria; Aquificae; Aquificales; Aquificaceae; Aquifex.
CC NCBI_TaxID=63363;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=VF5;
RC MEDLINE=98196666; PubMed=9537320;
RX
```

```
RA Deckert G., Warren P.V., Gaasterland T., Young W.G., Lenox A.L.,
RA Graham D.E., Overbeek R., Sneed M.A., Keller M., Aufay M., Huber R.,
RA Feldman R.A., Short J.M., Olson G.J., Swanson R.V.;
RT "The complete genome of the hyperthermophilic bacterium Aquifex
RT aeolicus.";
RL Nature 392:353-358(1998).
CC -1- SIMILARITY: SOME, TO R.PROWAZEKII RP189.
-----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
-----
DR EMBL; AE000723; AAC07153.1; -
DR PIR; B70395; B70395.
DR InterPro; IPR008921; Pol_clamp_load_C.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 350 AA; 40693 MW; 2949E786E1DAC2F9 CRC64;

Query Match      89.7%; Score 26; DB 1; Length 350;
Best Local Similarity 80.0%; Pred. No. 3.2e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 14 TKPPK 18

RESULT 34
TRMU_RICCN
ID TRMU_RICCN STANDARD; PRT; 370 AA.
AC Q9210;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Probable trna (5-methylaminomethyl-2-thiouridylate)-methyltransferase
DE (EC 2.1.1.61).
GN TRMU OR RC0410.
OS Rickettsia conorii.
CC Bacteria; Proteobacteria; Alphaproteobacteria; Rickettsiales;
CC Rickettsiaceae; Rickettsia.
CC NCBI_TaxID=781;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Malish 7;
RC MEDLINE=21442074; PubMed=11557893;
RA Ogata H., Audic S., Renesto-Audiffren P., Fournier P.-E., Barbe V.,
RA Sanson D., Roux V., Cossart P., Weissenbach J., Claverie J.-M.,
RA Raoult D.;
RT "Mechanisms of evolution in Rickettsia conorii and R. prowazekii.";
RL Science 293:2093-2098(2001).
CC -1- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + trna = S-adenosyl-L-
CC homocysteine + trna containing 5-methylaminomethyl-2-
CC thiouridylate.
CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -1- SIMILARITY: Belongs to the trmu family.
-----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
-----
DR EMBL; AE008605; AAL02948.1; -
DR PIR; B97751; B97751.
DR HAMAP; MF_00144; -; 1.
DR InterPro; IPR004506; TrmU.
DR Pfam; PF03054; trna_Me_trans; 1.
```

```
DR TIGRFAMS; TIGR00420; trmU; 1.
KW Transferase; Methyltransferase; tRNA processing; Complete proteome.
SQ SEQUENCE 370 AA; 40915 MW; 1884AC815E730CE3 CRC64;

Query Match      89.7%; Score 26; DB 1; Length 370;
Best Local Similarity 80.0%; Pred. No. 3.3e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
   |::|
Db 318 TKPPR 322

RESULT 35
RT31 HUMAN
ID RT31 HUMAN STANDARD; PRT; 395 AA.
AC Q92655; Q8WTV8;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE 28S ribosomal protein S31, mitochondrial precursor (S31mt) (MRP-S31)
DE (Imogen 38).
GN WRPS31 OR IMOGN38.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Skin;
RA Hutton J.C., Roep B.O.;
RA "Human Imogen 38. T-cell and antibody responses in newly diagnosed
RT diabetic subjects.";
RL Submitted (JAN-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Testis;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner K.H., Shenmen C.W., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Heide F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M.J., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Rana S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Rulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettaman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green B.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalhus D.B.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RN human and mouse cDNA sequences.";
RN Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [3]
RP IDENTIFICATION
RX MEDLINE=21276436; PubMed=11279123;
RA Koc E.C., Burkhardt W., Blackburn K., Moseley A., Spremulli L.L.;
RT "The small subunit of the mammalian mitochondrial ribosome:
RT identification of the full complement of ribosomal proteins present.";
RL J. Biol. Chem. 276:19363-19374(2001).
CC -1- SUBUNIT: Component of the mitochondrial ribosome small subunit
CC (28S) which comprises a 12S rRNA and about 30 distinct proteins.
CC -1- SUBCELLULAR LOCATION: Mitochondrial.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AP003000; BAB49856.1; ALT_INIT.
CC HAMAP; MF_00144; -; 1.
CC InterPro; IPR004506; TrmU.
CC Pfam; PF03054; tRNA_We_trans; 1.
CC TIGRFAMS; TIGR00420; trmU; 1.
KW Transferase; Methyltransferase; tRNA processing; Complete proteome.
SQ SEQUENCE 396 AA; 42537 MW; 66F118AA897E3086 CRC64;

CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; 268747; CAA92951.1; -.
CC EMBL; BC022045; AAH22045.1; -.
CC Genew; HGNC:16632; MRPS31.
CC GO; GO:0005739; C:mitochondrion; TAS.
CC GO; GO:0005739; C:mitochondrion; TAS.
KW Ribosomal protein; Mitochondrion; Transit peptide.
FT TRANSIT 1 65 MITOCHONDRION (POTENTIAL).
FT CHAIN 66 395 28S RIBOSOMAL PROTEIN S31.
FT CONFLICT 80 80 I -> T (IN REF. 1).
FT CONFLICT 132 132 A -> T (IN REF. 1).
FT CONFLICT 279 279 D -> N (IN REF. 1).
SQ SEQUENCE 395 AA; 45300 MW; E9410F46C94C6F3D CRC64;

Query Match      89.7%; Score 26; DB 1; Length 395;
Best Local Similarity 80.0%; Pred. No. 3.3e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
   |::|
Db 118 TKPPK 122

RESULT 36
TRMU RHIL0
ID TRMU RHIL0 STANDARD; PRT; 396 AA.
AC Q98H10;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Probable tRNA (5-methylaminomethyl-2-thiouridyate)-methyltransferase
DE (EC 2.1.1.61).
GN TRMU OR MLR2824.
OS Rhizobium loti (Mesorhizobium loti).
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Phyllobacteriaceae; Mesorhizobium.
OX NCBI_TaxID=381;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MAFP303099; PubMed=11214968;
RX MEDLINE=21082930; PubMed=11214968;
RA Kaneko T., Nakamura Y., Sato S., Asamizu E., Kato T., Sasamoto S.,
RA Watanabe A., Idegawa K., Ishikawa K., Kawashima K., Kimura T.,
RA Kishida Y., Kiyokawa C., Kohara M., Matsumoto M., Matsuno A.,
RA Mochizuki Y., Nakayama S., Nakazaki N., Shimpo S., Sugimoto M.,
RA Takeuchi C., Yamada M., Tabata S.;
RT "Complete genome structure of the nitrogen-fixing symbiotic bacterium
RL Mesorhizobium loti.";
RL DNA Res. 7:331-338(2000).
CC -1- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + tRNA = S-adenosyl-L-
CC homocysteine + tRNA containing 5-methylaminomethyl-2-
CC thiouridyate.
CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -1- SIMILARITY: Belongs to the trmU family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AP003000; BAB49856.1; ALT_INIT.
CC HAMAP; MF_00144; -; 1.
CC InterPro; IPR004506; TrmU.
CC Pfam; PF03054; tRNA_We_trans; 1.
CC TIGRFAMS; TIGR00420; trmU; 1.
KW Transferase; Methyltransferase; tRNA processing; Complete proteome.
SQ SEQUENCE 396 AA; 42537 MW; 66F118AA897E3086 CRC64;
```

```
Query Match      89.7%; Score 26; DB 1; Length 396;
Best Local Similarity 80.0%; Pred. No. 3.6e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 TKPPR 5
DB      320 TRPPR 324

RESULT 37
TRMU_BRUME
ID TRMU BRUME STANDARD; PRT; 398 AA.
AC Q8Y1L6;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DE Probable tRNA (5-methylaminomethyl-2-thiouridylate)-methyltransferase
DE (EC 2.1.1.61).
GN TRMU OR BME10428.
OS Brucella melitensis.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Brucellaceae; Brucella.
OX NCBI_TaxID=29459;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=16M / ATCC 23456 / Biotype 1;
RX MEDLINE=20020109; PubMed=1756886;
RA Delvecchio V.G., Kapatral V., Redkar R.J., Patra G., Mijer C., Los T.,
RA Ivanova N., Anderson I., Bhattacharya A., Lykidis A., Renik G.,
RA Jablonki L., Larsen L., P'Souza M., Bernal A., Mazur M., Goldsman E.,
RA Selkov E., Elzer P.H., Hagius S., O'Callaghan D., Letesson J.-J.,
RA Haselkorn R., Kyprides N., Overbeek R.;
RT "The genome sequence of the facultative intracellular pathogen
RT Brucella melitensis";
RL Proc. Natl. Acad. Sci. U.S.A. 99:443-448 (2002).
CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + tRNA = S-adenosyl-L-
CC thiouridylate.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the trmU family.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF009485; AAL51609.1; -
CC DR PIR; AF3305; AF3305.
CC DR HAMAP; MF_00144; -; 1.
CC InterPro; IPR004506; TrmU.
CC Pfam; PF03054; tRNA.Me.trans; 1.
CC TIGRFAMs; TIGR00420; trmU; 1.
CC TrnTransferase; Methyltransferase; tRNA processing; Complete proteome.
CC SEQUENCE 398 AA; 43175 MW; 93687D1D8B39ED3 CRC64;

Query Match      89.7%; Score 26; DB 1; Length 398;
Best Local Similarity 80.0%; Pred. No. 3.6e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 TKPPR 5
DB      322 TRPPR 326

RESULT 38
TRMU_BRUSU
ID TRMU BRUSU STANDARD; PRT; 398 AA.
AC Q8CY38;
DT 15-MAR-2004 (Rel. 43, Created)
DE Probable tRNA (5-methylaminomethyl-2-thiouridylate)-methyltransferase
DE (EC 2.1.1.61).
GN TRMU OR BME10428.
OS Brucella melitensis.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Brucellaceae; Brucella.
OX NCBI_TaxID=29459;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=16M / ATCC 23456 / Biotype 1;
RX MEDLINE=20020109; PubMed=1756886;
RA Delvecchio V.G., Kapatral V., Redkar R.J., Patra G., Mijer C., Los T.,
RA Ivanova N., Anderson I., Bhattacharya A., Lykidis A., Renik G.,
RA Jablonki L., Larsen L., P'Souza M., Bernal A., Mazur M., Goldsman E.,
RA Selkov E., Elzer P.H., Hagius S., O'Callaghan D., Letesson J.-J.,
RA Haselkorn R., Kyprides N., Overbeek R.;
RT "The genome sequence of the facultative intracellular pathogen
RT Brucella melitensis";
RL Proc. Natl. Acad. Sci. U.S.A. 99:443-448 (2002).
CC -!- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + tRNA = S-adenosyl-L-
CC thiouridylate.
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to the trmU family.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF009485; AAL51609.1; -
CC DR PIR; AF3305; AF3305.
CC DR HAMAP; MF_00144; -; 1.
CC InterPro; IPR004506; TrmU.
CC Pfam; PF03054; tRNA.Me.trans; 1.
CC TIGRFAMs; TIGR00420; trmU; 1.
CC TrnTransferase; Methyltransferase; tRNA processing; Complete proteome.
CC SEQUENCE 398 AA; 43175 MW; 93687D1D8B39ED3 CRC64;

Query Match      89.7%; Score 26; DB 1; Length 398;
Best Local Similarity 80.0%; Pred. No. 3.6e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 TKPPR 5
DB      322 TRPPR 326

RESULT 39
VGLD_HSVEA
ID VGLD HSVEA STANDARD; PRT; 402 AA.
AC P24872;
DT 01-MAR-1992 (Rel. 21, Created)
DT 01-AUG-1992 (Rel. 23, Last sequence update)
DE Probable tRNA (5-methylaminomethyl-2-thiouridylate)-methyltransferase
DE (EC 2.1.1.61).
GN VGLD OR G17/18 OR 72.
OS Equine herpesvirus type 1 (strain ABI) (EHV-1).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
OX NCBI_TaxID=10328;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE=92268882; PubMed=1316942;
RX Elton D.M., Halliburton I.W., Killington R.A., Meredith D.M.,
RA Bonas W.A.;
RT "Identification of the equine herpesvirus type 1 glycoprotein 17/18
RT as a homologue of herpes simplex virus glycoprotein D.";
RT
```

J. Gen. Virol. 73:1227-1233(1992).

[2]
RN SEQUENCE OF 242-402 FROM N.A.
RP MEDLINE=91276272; PubMed=1647359;
RX Elton D.W., Halliburton I.W., Killington R.A., Meredith D.M.,
RA Bonas W.A.;
RT "Sequence analysis of the 4.7-Kb BamHI-EcoRI fragment of the equine
RT herpesvirus type-1 short unique region.";
RL Gen 101:203-208(1991).
RN -1- SIMILARITY: Belongs to the herpesviruses glycoprotein D family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).

EMBL; M60946; AAA46087.1; -
EMBL; M36299; AAA65546.1; -
DR InterPro; IPR002896; Herpes glycop D.
DR InterPro; IPR007110; Ig-like.
DR Pfam; PF01537; Herpes Glycop D; 1.
KW Glycoprotein; Transmembrane; Signal.
FT SIGNAL 1 30 POTENTIAL.
FT CHAIN 31 402 GLYCOPROTEIN D.
FT DOMAIN 31 355 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 356 372 POTENTIAL.
FT DOMAIN 373 402 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 53 53 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 61 61 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 297 297 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 346 346 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 402 AA; 45211 MW; 78A0593232D0238C CRC64;

Query Match 89.7%; Score 26; DB 1; Length 402;
Best Local Similarity 80.0%; Pred. No. 3.6e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 337 TKPPK 341

RESULT 40
ID_CCKR_HUMAN STANDARD; PRT; 428 AA.
AC P32238;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-OCT-1993 (Rel. 27, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Cholecystokinin type A receptor (CCK-A receptor) (CCK-AR).
GN CCKAR OR CCKRA.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
OX NCBI_TaxID=9606;
RN [1]
RN SEQUENCE FROM N.A.
RP TISSUE=Gall bladder;
RX MEDLINE=93277532; PubMed=8503909;
RA Ulrich C.D., Ferber I., Holicky E., Hadac E., Buell G.,
RA Miller L.J.;
RT "Molecular cloning and functional expression of the human gallbladder
RT cholecystokinin A receptor.";
RL Biochem. Biophys. Res. Commun. 193:204-211(1993).
RN [2]
RN SEQUENCE FROM N.A.
RP MEDLINE=93343941; PubMed=8343165;
RX Wank S.A., de Weerth A., Pilegna J.R., Huppi K.;
RT "Molecular cloning, functional expression and chromosomal
RT localization of the human cholecystokinin type A receptor.";

Biochem. Biophys. Res. Commun. 194:811-818(1993).

[3]
RN SEQUENCE FROM N.A.
RP MEDLINE=96029343; PubMed=7557108;
RX Miller L.J., Holicky E.L., Ulrich C.D., Wieben E.D.;
RA "Abnormal processing of the human cholecystokinin receptor gene in
RT association with gallstones and obesity.";
RL Gastroenterology 109:1375-1380(1995).
RN [4]
RN SEQUENCE FROM N.A.
RP TISSUE=Peripheral blood leukocytes;
RX MEDLINE=20145045; PubMed=10682840;
RA Funakoshi A., Miyasaka K., Matsumoto H., Yamamori S., Takiguchi S.,
RA Katsuka K., Takata Y., Matsue K., Kono A., Shimokata H.;
RT "Gene structure of human cholecystokinin (CCK) type-A receptor: body
RT fat content is related to CCK type-A receptor gene promoter
RT polymorphism.";
RL FEBS Lett. 466:264-266(2000).
RN [5]
RN SEQUENCE FROM N.A.
RP TISSUE=Stomach;
RX Kopatz S.A., Aronstam R.S., Sharma S.V.;
RT "cDNA clones of human proteins involved in signal transduction
RT signalled by the Guthrie cDNA resource center (www.cdna.org).";
RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Receptor for cholecystokinin. Has a 1000 fold affinity
CC for CCK rather than for gastrin. It modulates feeding and
CC dopamine-induced behavior in the central and peripheral nervous
CC system. This receptor mediates its action by association with G
CC proteins that activate a phosphatidylinositol-calcium second
CC messenger system.
CC -!- SUBCELLULAR LOCATION: Integral membrane protein.
CC -!- SIMILARITY: Belongs to family 1 of G-protein coupled receptors.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).

EMBL; L13605; AAA35659.1; -
EMBL; L19315; AAA02819.1; -
EMBL; U23430; AAA91123.1; -
EMBL; U23427; AAA91123.1; JOINED.
EMBL; U23428; AAA91123.1; JOINED.
EMBL; U23429; AAA91123.1; JOINED.
EMBL; D85606; BAA90879.1; -
EMBL; AY322549; AAP84362.1; -
DR FIR; JN0692; JN0692.
DR PDB; 1D6G; 17-NOV-99.
DR PDB; 1H2N; 25-APR-01.
DR Genew; HGNC:1570; CCKAR.
DR MIM; 118444;
DR GO; GO:0008887; C:integral to plasma membrane; TAS.
DR GO; GO:0004951; F:cholecystokinin receptor activity; TAS.
DR GO; GO:0007204; P:cytosolic calcium ion concentration elevation; TAS.
DR GO; GO:0007586; P:digestion; TAS.
DR GO; GO:0007631; P:feeding behavior; TAS.
DR GO; GO:0007584; P:response to nutrients; TAS.
DR InterPro; IPR000276; GPCR_Rhodpsn.
DR Pfam; PF00001; 7tm1; 1.
DR PRINTS; PR00237; GPCRHHODPSN.
DR PROSITE; PS00237; G_PROTEIN_RECF1_1; 1.
DR PROSITE; PS0262; G_PROTEIN_RECF1_2; 1.
KW G-protein coupled receptor; Transmembrane; Glycoprotein; Lipoprotein;
KW Palmitate; 3D-structure.
FT DOMAIN 1 41 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 42 67 1 (POTENTIAL)
FT DOMAIN 68 77 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 78 104 2 (POTENTIAL).
FT DOMAIN 105 115 EXTRACELLULAR (POTENTIAL).

FT TRANSMEM 116 137 3 (POTENTIAL).
 FT DOMAIN 138 157 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 178 178 4 (POTENTIAL).
 FT DOMAIN 179 210 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 211 234 5 (POTENTIAL).
 FT DOMAIN 235 313 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 314 334 6 (POTENTIAL).
 FT DOMAIN 335 349 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 350 373 7 (POTENTIAL).
 FT DOMAIN 374 428 CYTOPLASMIC (POTENTIAL).
 FT CARBOHYD 10 10 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 24 24 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 190 190 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT DISULFID 114 196 BY SIMILARITY.
 FT LIPID 387 S-palmitoyl cysteine (By similarity).
 SQ SEQUENCE 428 AA; 47841 MW; A688FABDA05610 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 428;
 Best Local Similarity 80.0%; Pred. No. 3.9e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
 :|||
 Db 274 TRPPR 278

RESULT 41
 NOCT_MOUSE STANDARD; PRT; 429 AA.
 AC Q35710; Q9Q2G9;
 DT 15-JUL-1999 (Rel. 38, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Nocturnin (CCR4 protein homolog).
 GN CCRN4L OR NOC OR CCR4.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BALB/c; TISSUE=Brain;
 RX MEDLINE=99453012; PubMed=10521507;
 RA Dupressoir A., Barbot W., Loireau M.P., Heidmann T.;
 RT "Characterization of a mammalian gene related to the yeast CCR4
 general transcription factor and revealed by transposon insertion.";
 RL J. Biol. Chem. 274:31068-31075(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BALB/c; TISSUE=Retina;
 RX MEDLINE=22944738; PubMed=11394964;
 RA Wang Y., Osterbur D.L., Megaw P.L., Tosini G., Fukuhara C.,
 Green C.B., Besharse J.C.;
 RT "Rhythmic expression of Nocturnin mRNA in multiple tissues of the
 mouse.";
 RL BMC Dev. Biol. 1:9-9(2001).
 RN [3]
 RP SEQUENCE OF 62-429 FROM N.A.
 RC STRAIN=DBA/2J; TISSUE=Liver;
 RX MEDLINE=97190339; PubMed=9038221;
 RA Puech A., Dupressoir A., Loireau M.P., Mattei M.-G., Heidmann T.;
 RT "Characterization of two age-induced intracisternal A-particle-related
 transcripts in the mouse liver. Transcriptional read-through into an
 open reading frame with similarities to the yeast ccr4 transcription
 factor.";
 RL J. Biol. Chem. 272:5995-6003(1997).
 CC -!- FUNCTION: Component of the circadian clock or downstream effector
 of clock function. Exhibits a high amplitude circadian rhythm with
 maximal levels in early evening. In constant darkness or constant
 light, the amplitude of the rhythm decreases (by similarity).
 CC -!- SIMILARITY: Belongs to the CCR4/nocturnin family.
 CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration

CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; AF183960; AAD56547.1; -;
 DR EMBL; AF199491; AAG01384.1; -;
 DR EMBL; U70139; AAB62717.1; ALT_FRAME.
 DR MGD; MGI:109382; Ccrn4l.
 DR InterPro; IPR005135; Exo_endo_phos.
 DR Pfam; PF03372; Exo_endo_phos; 1.
 KW Biological rhythms.
 FT CONFLICT 123 125 YOR -> LPA (IN REF. 3).
 SQ SEQUENCE 429 AA; 48300 MW; CB9FBE5D84E13942 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 429;
 Best Local Similarity 80.0%; Pred. No. 3.9e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
 :|||
 Db 118 TRPPR 122

RESULT 42
 NOCT_HUMAN STANDARD; PRT; 431 AA.
 AC Q9UK39; Q9HD93; Q9HD94; Q9HD95;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Nocturnin (CCR4 protein homolog).
 GN CCRN4L OR NOC OR CCR4.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC MEDLINE=99453012; PubMed=10521507;
 RA Dupressoir A., Barbot W., Loireau M.P., Heidmann T.;
 RT "Characterization of a mammalian gene related to the yeast CCR4
 general transcription factor and revealed by transposon insertion.";
 RL J. Biol. Chem. 274:31068-31075(1999).
 RN [2]
 RP SEQUENCE OF 67-431 FROM N.A.
 RA Wang Y., Osterbur D.L., Green C.B., Besharse J.C.;
 RT "Mammalian homologs of Xenopus nocturnin: conservation of structure
 and circadian regulation.";
 RL Submitted (OCT-1999) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: Component of the circadian clock or downstream effector
 of clock function. Exhibits a high amplitude circadian rhythm with
 maximal levels in early evening. In constant darkness or constant
 light, the amplitude of the rhythm decreases (by similarity).
 CC -!- SIMILARITY: Belongs to the CCR4/nocturnin family.
 CC -----
 CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; AF183961; AAD56548.1; -;
 DR EMBL; AF199492; AAG01387.1; -;
 DR EMBL; AF199493; AAG01388.1; -;
 DR EMBL; AF199494; AAG01389.1; -;
 DR Genew; HGNC:14254; CCRN4L.
 DR GO; GO:0005634; Cinnuleus; TAS.
 DR GO; GO:0003700; F:transcription factor activity; TAS.

DR GO: GO:0006366; P:transcription from Pol II promoter; TAS.
DR InterPro: IPR005135; Exo_endo_phos.
DR Pfam: PF03372; Exo_endo_phos; 1.
KW Biological rhythms. 69 T -> N (IN REF. 2).
FT CONFLICT 69 G -> A (IN REF. 2).
FT CONFLICT 77 A -> T (IN REF. 2); AAG01389).
FT CONFLICT 266 N -> S (IN REF. 2).
FT CONFLICT 341 N -> S (IN REF. 2).
SQ SEQUENCE 431 AA; 48150 MW; B61EF484EBD29AF5 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 431;
Best Local Similarity 80.0%; Pred. No. 3.9e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 120 TRPPR 124

RESULT 43

ENGA_CLOPE STANDARD; PRT; 438 AA.
AC Q8XJ1;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE GTP-binding protein engA.
GN ENGA OR CPEI755.
OS Clostridium perfringens.
OC Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;
OC Clostridium.
OX NCBI_TaxID=1502;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=13 / Type A;
RX MEDLINE=21664373; PubMed=11792842;
RA Shimizu T., Ohtani K., Hirakawa H., Ohshima K., Yamashita A.,
RA Shiba T., Ogasawara N., Hattori M., Kuhara S., Hayashi H.;
RT "Complete genome sequence of Clostridium perfringens, an anaerobic
flesh-eater";
RL Proc. Natl. Acad. Sci. U.S.A. 99:996-1001(2002).

CC -!- FUNCTION: GTPase of unknown physiological role.
CC -!- SIMILARITY: Belongs to the era/trmE family of GTP-binding
proteins. EngA subfamily.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
between the Swiss Institute of Bioinformatics and the EMBL outstation -
the European Bioinformatics Institute. There are no restrictions on its
use by non-profit institutions as long as its content is in no way
modified and this statement is not removed. Usage by and for commercial
entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
or send an email to license@isb-sib.ch).

EMBL: AP003191; BAB81461.1; -

DR HAMAP, MF_00195; 1.
DR InterPro: IPR003593; AAA_ATPase.
DR InterPro: IPR005289; GTP-Binding_dom.
DR InterPro: IPR006073; GTP1_OBG.
DR InterPro: IPR002917; MMR_HSR1.
DR InterPro: IPR01806; Ras_transfmg.
DR InterPro: IPR005225; Small_GTP.
DR Pfam: PF01926; MMR_HSR1; 1.
DR PRINTS: PR00326; GTP1_OBG.
DR PRINTS: PR00449; RASTNSFRMG.
DR SMART: SM00382; AAA; 2.
DR TIGRFAMs: TIGR00650; MG442; 2.
DR TIGRFAMs: TIGR00231; small_GTP; 2.
KW GTP-binding; Repeat; Complete proteome.
FT NP_BIND 10 17 GTP 1 (POTENTIAL).
FT NP_BIND 57 61 GTP 1 (POTENTIAL).
FT NP_BIND 120 123 GTP 1 (POTENTIAL).
FT NP_BIND 183 190 GTP 2 (POTENTIAL).
FT NP_BIND 230 234 GTP 2 (POTENTIAL).

FT NP_BIND 295 298 GTP 2 (POTENTIAL).
SQ SEQUENCE 438 AA; 49586 MW; 9658B0BC98398396 CRC64;
Query Match 89.7%; Score 26; DB 1; Length 438;
Best Local Similarity 80.0%; Pred. No. 4e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
Db 391 TKPPK 395

RESULT 44

SRMB_HAEIN STANDARD; PRT; 439 AA.
AC P44701;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE ATP-dependent RNA helicase srmb homolog.
GN SRMB OR H10422.
OS Haemophilus influenzae.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
OC Pasteurellaceae; Haemophilus.
OX NCBI_TaxID=727;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=rd / KW20 / ATCC 51907;
RX MEDLINE=95350630; PubMed=7542800;
RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
RA Kerlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
RA McKeeney K., Sutton R., Fitzhugh W., Fields C.A., Gocayne J.D.,
RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
RA Uettermann T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,
RA Fine L.D., Fritchman J.L., Fuhrmann J.L., Geoghagen N.S.M.,
RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
RA Venter J.C.;
RT "Whole-genome random sequencing and assembly of Haemophilus influenzae
Rd";
RL Science 269:496-512(1995).
CC -!- FUNCTION: RNA-DEPENDENT ATPASE ACTIVITY. PROBABLY INTERACTS
WITH 23S RIBOSOMAL RNA (BY SIMILARITY).
CC -!- SIMILARITY: Belongs to the DEAD box helicase family.

CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
between the Swiss Institute of Bioinformatics and the EMBL outstation -
the European Bioinformatics Institute. There are no restrictions on its
use by non-profit institutions as long as its content is in no way
modified and this statement is not removed. Usage by and for commercial
entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
or send an email to license@isb-sib.ch).

EMBL: U32725; AAC22078.1; -

DR PIR: H64066; H64066.
DR HSSP: Q58083; 1HV8.
DR TIGR: H10422;
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR000629; DEAD_box.
DR InterPro: IPR001650; Helicase_C.
DR Pfam: PF00270; DEAD; 1.
DR Pfam: PF00271; helicase_C; 1.
DR SMART: SM00487; DEXDC; 1.
DR SMART: SM00490; HELIC; 1.
DR PROSITE: PS00039; DEAD_ATP_HELICASE; 1.
KW Helicase; ATP-binding; RNA-binding; Complete proteome.
FT NP_BIND 48 55 ATP (POTENTIAL).
FT SITE 157 160 DEAD BOX.
SQ SEQUENCE 439 AA; 49805 MW; 7543942CE35B20C4 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 439;
Best Local Similarity 80.0%; Pred. No. 4e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPR 5
Db 383 TRPK 387

RESULT 45
VGLD_HSVK STANDARD; PRT; 442 AA.

ID VGLD_HSVK STANDARD; PRT; 442 AA.
AC P22454;
DT 01-AUG-1991 (Rel. 19, Created)
DT 01-AUG-1991 (Rel. 19, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Glycoprotein D precursor (Glycoprotein 17/18).
GN GD OR GP17/18 OR 72.
OS Equine herpesvirus type 1 (strain Kentucky A) (EHV-1).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
OC NCBI_TaxID=10329;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91082407; PubMed=1845821;
RA "Flowers C.C., Eastman E.M., O'Callaghan D.J.;
RT "Sequence analysis of a glycoprotein D gene homolog within the unique
RT short segment of the EHV-1 genome.";
RL Virology 180:175-184 (1991).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=92263758; PubMed=1316573;
RA "Colle C.F. III, Flowers C.C., O'Callaghan D.J.;
RT "Open reading frames encoding a protein kinase, homolog of
RT glycoprotein gx of pseudorabies virus, and a novel glycoprotein map
RT within the unique short segment of equine herpesvirus type 1.";
RL Virology 188:545-557 (1992).
CC -1- SIMILARITY: Belongs to the herpesviruses glycoprotein D family.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M62923; AAA46081.1; -;
DR EMBL; M66931; -; NOT ANNOTATED CDS.
DR PIR; A38518; VGBEEA.
DR InterPro; IPR002896; Herpes glycop_D.
DR Pfam; PF01537; Herpes glycop_D; 1.
KW Glycoprotein; Signal; Transmembrane.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 442 GLYCOPROTEIN D.
FT DOMAIN 20 405 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 406 422 POTENTIAL.
FT DOMAIN 423 442 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 103 103 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 111 111 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 347 347 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 396 396 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 442 AA; 49908 MW; 332CDCA9C9762F05 CRC64;

Query Match 89.7%; Score 26; DB 1; Length 442;
Best Local Similarity 80.0%; Pred. No. 4e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPR 5
Db 387 TRPK 391

RESULT 46
VGLD_HSVK STANDARD; PRT; 463 AA.

ID ENGA_BIFLO STANDARD; PRT; 463 AA.
AC Q8G6J8;
DT 15-MAR-2004 (Rel. 43, Created)
DT 15-MAR-2004 (Rel. 43, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)

VGLD_HSVB STANDARD; PRT; 452 AA.

ID VGLD_HSVB STANDARD; PRT; 452 AA.
AC P24379;
DT 01-MAR-1992 (Rel. 21, Created)
DT 01-MAR-1992 (Rel. 21, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Glycoprotein D precursor (Glycoprotein 17/18).
GN GD OR GP17/18 OR 72.
OS Equine herpesvirus type 1 (strain Ab4p) (EHV-1), and
OS Equine herpesvirus type 1 (strain Kentucky D) (EHV-1).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
OC NCBI_TaxID=31520; 10330;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=AB4P;
RX MEDLINE=92295566; PubMed=1318606;
RA Telford E.A.R., Watson M.S., McBride K., Davison A.J.;
RT "The DNA sequence of equine herpesvirus-1.";
RL Virology 189:304-316 (1992).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=Kentucky D;
RX MEDLINE=91108393; PubMed=2177089;
RA "Adonnet J.-C., Winslow J., Allen G., Paoletti E.;
RT "Equine herpesvirus type 1 unique short fragment encodes
RT glycoproteins with homology to herpes simplex virus type 1 GD, GI and
RT GE.";
RL J. Gen. Virol. 71:2969-2978 (1990).
CC -1- SIMILARITY: Belongs to the herpesviruses glycoprotein D family.
CC -1- CAUTION: It is uncertain whether Met-1 or Met-51 is the initiator.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M86664; AAB02507.1; -;
DR PIR; I36802; VGBEG3.
DR InterPro; IPR002896; Herpes glycop_D.
DR PIR; I36802; VGBEG3.
DR InterPro; IPR007110; Ig-like_D; 1.
DR Pfam; PF01537; Herpes glycop_D; 1.
KW Glycoprotein; Signal; Transmembrane.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 452 GLYCOPROTEIN D.
FT DOMAIN 20 405 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 406 422 POTENTIAL.
FT DOMAIN 423 452 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 103 103 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 111 111 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 347 347 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 396 396 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 452 AA; 51099 MW; CFS1E914F7F2E9DC CRC64;

Query Match 89.7%; Score 26; DB 1; Length 452;
Best Local Similarity 80.0%; Pred. No. 4.1e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPR 5
Db 387 TRPK 391

RESULT 47
ENGA_BIFLO STANDARD; PRT; 463 AA.

ID ENGA_BIFLO STANDARD; PRT; 463 AA.
AC Q8G6J8;
DT 15-MAR-2004 (Rel. 43, Created)
DT 15-MAR-2004 (Rel. 43, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)

DE GTP-binding protein engA.
GN ENGA OR BL0738.
OS Bifidobacterium longum.
CC Bacteria; Actinobacteridae; Actinobacteriales; Bifidobacteriales;
CC Bifidobacteriaceae; Bifidobacterium.
OX NCBI_TaxID=216816;
RN [1]_TaxID=216816;
RP SEQUENCE FROM N.A.
RC STRAIN=NCC 2705;
RX MEDLINE=2294977; PubMed=13381787;
RA Scheil M.A., Karmirantzou M., Snel B., Vilanova D., Berger B.,
RA Pessi G., Zwaan H.M.-C., Desiere F., Bork P., Delley M.,
RA Pridmore R.D., Arigoni F.,
RT "The genome sequence of Bifidobacterium longum reflects its adaptation
RT to the human gastrointestinal tract".
RL Proc. Natl. Acad. Sci. U.S.A. 99:14422-14427(2002).
CC -!- FUNCTION: GTPase of unknown physiological role.
CC -!- SIMILARITY: Belongs to the era/trmE family of GTP-binding
CC proteins. EngA subfamily.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
CC ENBL; AEO14696; AAN24555.1; -.
CC HAMAP; MF_00195; 1.
CC InterPro; IPR003593; AAA ATPase.
CC InterPro; IPR005289; GTP-binding_dom.
CC InterPro; IPR006073; GTP1_OBG.
CC InterPro; IPR002917; MMR HSR1.
CC InterPro; IPR005225; Small GTP.
CC Pfam; PF01926; MMR HSR1. 1.
CC PRINTS; PR00326; GTP1_OBG.
CC SMART; SMO0382; AAA; 2.
CC TIGRFAMs; TIGR00650; MG442; 2.
CC TIGRFAMs; TIGR00231; small_GTP; 2.
CC TIGRFAMs; Repeat; Complete proteome.
CC KW NP-BIND; 33 40 GTP 1 (POTENTIAL).
CC FT NP-BIND 80 84 GTP 1 (POTENTIAL).
CC FT NP-BIND 142 145 GTP 1 (POTENTIAL).
CC FT NP-BIND 208 215 GTP 2 (POTENTIAL).
CC FT NP-BIND 255 259 GTP 2 (POTENTIAL).
CC FT NP-BIND 320 323 GTP 2 (POTENTIAL).
CC SQ SEQUENCE 463 AA; 51186 MW; 7F986F8E041367AE CRC64;

Query Match 89.7%; Score 26; DB 1; Length 463;
Best Local Similarity 80.0%; Pred. No. 4.2e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPRP 5
Db 414 TRPRP 418

RESULT 48
EM55 HUMAN STANDARD; PRT; 466 AA.
AC Q00013;
DT 01-DEC-1992 (Rel. 24, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE 55 kDa erythrocyte membrane protein (p55) (Membrane protein,
DE palmitoylated 1).
GN MPPI OR EMP55.
OS Homo sapiens (Human).
CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
CC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]

RP SEQUENCE FROM N.A. AND PARTIAL SEQUENCE.
RC TISSUE=Reticulocytes;
RX MEDLINE=91319732; PubMed=1713685;
RA Ruff P., Speicher D.W., Husain-Chishti A.;
RT "Molecular identification of a major palmitoylated erythrocyte
RT membrane protein containing the src homology 3 motif".
RL Proc. Natl. Acad. Sci. U.S.A. 88:6595-6599(1991).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=93244792; PubMed=1301163;
RA Metznerberg A.B., Gitschler J.;
RT "The gene encoding the palmitoylated erythrocyte membrane protein,
RT p55, originates at the CpG island 3' to the factor VIII gene".
RL Hum. Mol. Genet. 1:97-101(1992).
RN [3]
RP SEQUENCE FROM N.A.
RC TISSUE=Lung;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.D., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.A., Loquellano N.A., Peters G.J., Abramson R.D., Mullady S.J.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettelman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smallos D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences".
CC Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -!- SUBUNIT: Interacts with DLG5.
CC -!- SUBCELLULAR LOCATION: Membrane-associated.
CC -!- PTM: Extensively palmitoylated.
CC -!- SIMILARITY: Belongs to the MAGUK family.
CC -!- SIMILARITY: Contains 1 PDZ/DHR domain.
CC -!- SIMILARITY: Contains 1 SH3 domain.
CC -!- SIMILARITY: Contains 1 guanylate kinase-like domain.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
CC ENBL; M64925; AAA60059.1; -.
CC EMBL; M87059; AAA60060.1; -.
CC EMBL; U39611; AAD14835.1; -.
CC EMBL; BC002392; AAH02392.1; -.
CC HSP; O14936; 1KWA.
CC Genew; HGNC:7219; MPPI.
CC MIM; 305360; -.
CC GO; GO:0005887; C:integral to plasma membrane; TAS.
CC GO; GO:0005624; C:membrane fraction; TAS.
CC GO; GO:0004385; F:guanylate kinase activity; TAS.
CC GO; GO:0007165; P:signal transduction; TAS.
CC InterPro; IPR008144; Guanylate_kin.
CC InterPro; IPR001478; PDZ.
CC InterPro; IPR001452; SH3.
CC Pfam; PF00625; Guanylate_kin; 1.
CC Pfam; PF00595; PDZ; 1.
CC Pfam; PF00018; SH3; 1.
CC ProDom; PD000066; SH3; 1.

```

DR SMART; SM00072; GUKC; 1.
DR SMART; SM00228; PDZ; 1.
DR SMART; SM00326; SH3; 1.
DR PROSITE; PS00856; GUANYLATE KINASE 1; 1.
DR PROSITE; PS00052; GUANYLATE_KINASE_2; 1.
DR PROSITE; PS0106; PDZ; 1.
DR PROSITE; PS0002; SH3; 1.
KW Membrane, Erythrocyte; SH3 domain; Lipoprotein; Palmitate;
KW Polymorphism.
FT DOMAIN 71 152 PDZ.
FT DOMAIN 158 228 SH3.
FT DOMAIN 280 466 GUANYLATE KINASE.
FT VARIANT 448 448 E -> Q (in dbSNP:14092).
FT FTID=VAR_011914.
SQ SEQUENCE 466 AA; 52296 MW; DC68AA68F49A26E CRC64;

Query Match 89.7%; Score 26; DB 1; Length 466;
Best Local Similarity 80.0%; Pred. No. 4.2e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 318 TRPPR 322

RESULT 49
ENP6_HUMAN STANDARD; PRT; 484 AA.
ID ENP6_HUMAN STANDARD; PRT; 484 AA.
AC O75354; Q9UUD1;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Ectonucleoside triphosphate diphosphohydrolase 6 (EC 3.6.1.16)
DE (NTPases) (CD39 antigen-like 2).
DE ENTPD6 OR CD39L2.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
CX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Keratinocytes;
RX MEDLINE=98341119; PubMed=9676430;
RA Chadwick B.P., Frischauf A.-M.;
RT "The CD39-like gene family: identification of three new human members
RT (CD39L2, CD39L3, and CD39L4), their murine homologues, and a member of
RT the gene family from Drosophila melanogaster.";
RL Genomics 50:357-367(1998).
[2]
RP SEQUENCE FROM N.A.
RX MEDLINE=21638749; PubMed=11780052;
RA Deloukas P., Matthews L.H., Ashurst J., Burton J., Gilbert J.G.R.,
RA Jones M., Stavrides G., Almeida J.P., Babbage A.K., Bagguley C.L.,
RA Bailey J., Barlow K.F., Bates K.N., Beard L.M., Beare D.M.,
RA Beasley O.P., Bird C.P., Blakey S.E., Bridgeman A.M., Brown A.J.,
RA Buck D., Burrill W.D., Butler A.P., Carder C., Carter N.P.,
RA Chapman J.C., Clamp M., Clark G., Clark L.N., Clark S.Y., Clee C.M.,
RA Clegg S., Cobley V.E., Collier R.E., Connor R.E., Corby N.R.,
RA Coulson A., Coville G.J., Deadman R., Dami P.D., Dunn M.,
RA Ellington A.G., Frankland J.A., Fraser A., French L., Garner P.,
RA Grainger D.V., Griffiths C., Griffiths M.N.D., Gwilliam R., Hall R.E.,
RA Hammond S., Harley J.L., Heath P.D., Ho S., Holden J.L., Howden P.J.,
RA Huckle E., Hunt A.R., Hunt S.E., Jekosch K., Johnson C.M., Johnson D.,
RA Kay M.P., Kimberley A.M., King A., Knights A., Laird G.K., Lawlor S.,
RA Lehaevaisho M.H., Leverisa M.C., Lloyd C., Lloyd D.M., Lovell J.D.,
RA Marsh V.L., Martin S.L., McConachie L.J., McLeay K., McMurray A.A.,
RA Milne S.A., Mistry D., Moore M.J.F., Mullikin J.C., Nickerson T.,
RA Oliver K., Parker A., Patel R., Pearce T.A.V., Peck A.I.,
RA Phillimore B.J.C.T., Pratchalingam S.R., Plumb R.W., Ramsay H.,
RA Rice C.D., Ross M.P., Scott C.E., Sehra H.K., Showkhen R., Sims S.,
RA Skuce C.D., Smith M.L., Soderlund C., Steward C.A., Sulston J.E.,
RA Swann R.M., Sycamore N., Taylor R., Tee L., Thomas D.W., Thorpe A.,
RA Tracey A., Tromans A.C., Vaudin M., Wall M., Wallis J.M.,

```

```

RA Whitehead S.L., Whittaker P., Willey D.L., Williams L., Williams S.A.,
RA Wilming L., Wray P.W., Hubbard T., Durbin R.M., Bentley D.R., Beck S.,
RA Rogers J.;
RT "The DNA sequence and comparative analysis of human chromosome 20.";
RL Nature 414:865-871(2001).
CC -1- FUNCTION: Might support glycosylation reactions in the Golgi
CC apparatus and when released from cells, might catalyze the
CC hydrolysis of extracellular nucleotides. Hydrolyzes preferentially
CC nucleoside 5'-diphosphates, nucleoside 5'-triphosphates are
CC hydrolyzed only to a minor extent, there is no hydrolysis of
CC nucleoside 5'-monophosphates. The order of activity with different
CC substrates is GDP > IDP > UDP = CDP >> ADP (By similarity).
CC -1- CATALYTIC ACTIVITY: A nucleoside diphosphate + H(2)O = a
CC nucleotide + phosphate.
CC -1- COFACTOR: Requires calcium and magnesium (By similarity).
CC -1- SUBCELLULAR LOCATION: TYPE II MEMBRANE PROTEIN. GOLGI. BUT ALSO
CC OCCURS IN A SOLUBLE EXTRACELLULAR FORM (BY SIMILARITY).
CC -1- TISSUE SPECIFICITY: Expressed in most tissues.
CC -1- SIMILARITY: Belongs to the GDAL / CD39 NTPase family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AF039916; AAC39883.1; -.
CC EMBL; AL035252; CAB41571.1; -.
CC Genbank; HGNC:3368; ENTPD6.
CC MIM; 603160; -.
CC InterPro; IPR000407; GDAL_CD39_NTPase.
CC Pfam; PF01150; GDAL_CD39_1.
CC PROSITE; PS01238; GDAL_CD39_NTPase; FALSE NEG.
KW Hydrolyase; Transmembrane; Glycoprotein; Calcium; Magnesium;
KW Signal-anchor; Go-gi stack.
FT DOMAIN 1 39
FT TRANSMEM 40 60
FT CYTOPLASMIC (POTENTIAL).
FT SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
FT (POTENTIAL).
FT LUMENAL (POTENTIAL).
FT N-LINKED (GLCNAc...) (POTENTIAL).
FT CARBOHYD 220 220 N-LINKED (GLCNAc...) (POTENTIAL).
FT CARBOHYD 284 284 V -> L (IN REF. 2).
FT CONFLICT 138 138 E -> K (IN REF. 2).
FT CONFLICT 202 202
FT SEQUENCE 484 AA; 53233 MW; 27334E29DDB8D64C CRC64;

Query Match 89.7%; Score 26; DB 1; Length 484;
Best Local Similarity 80.0%; Pred. No. 4.4e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 121 TRPPR 125

RESULT 50
AMEA_AERPE STANDARD; PRT; 524 AA.
AC Q9Y935;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Probable cytosol aminopeptidase (EC 3.4.11.1) (Leucine aminopeptidase)
DE (LAP) (Leucyl aminopeptidase).
DE PEPA OR APE2450.
GN Aeropyrum pernix.
OS Archaea; Crenarchaeota; Thermoprotei; Desulfurococcaceae;
OC Desulfurococcaceae; Aeropyrum.
OX NCBI_TaxID=56636;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=K1;

```

PX MEDLINE-99310339; PubMed-10382966;
RA Kawarabayashi Y., Hino Y., Horikawa H., Yamazaki S., Hatakeyama Y.,
RA Jinno K., Takahashi M., Sekine M., Baba S.-I., Ankai A., Kosugi H.,
RA Hosoyama A., Fukui S., Nagai Y., Nishijima K., Nakazawa H.,
RA Takamiya M., Masuda S., Funahashi T., Tanaka T., Kudo Y.,
RA Yamazaki J., Kishida N., Oguchi A., Aoki K.-I., Kubota K.,
RA Nakamura Y., Nomura N., Sako Y., Kikuchi H.,
RT "Complete genome sequence of an aerobic hyper-thermophilic
RT crenarchaeon, Aeropyrum pernix K1.";
RL DNA Res. 6:83-101(1999).
CC -!- FUNCTION: Presumably involved in the processing and regular
CC turnover of intracellular proteins. Catalyzes the removal of
CC unsubstituted N-terminal amino acids from various peptides (By
CC similarity).
CC -!- CATALYTIC ACTIVITY: Release of an N-terminal amino acid, Xaa-|-
CC including Pro although not Arg or Lys, and Xbb may be Pro.
CC -!- COFACTOR: Binds 2 manganese ions per subunit (By similarity).
CC -!- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC -!- SIMILARITY: Belongs to peptidase family M17.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: AP000064; BAA81465.1; -;
DR PIR: A72476; A72476.
DR HSP: P00727; ILAP.
DR MEROPS: M17.0PW; -;
DR HAMAP: MF 00181; -; 1.
DR InterPro: IPR000819; Peptidase M17_C.
DR InterPro: IPR008283; Peptidase M17_N.
DR Pfam: PF00883; Peptidase M17; I.
DR Pfam: PF02789; Peptidase M17_N; 1.
DR PRINTS: PR00481; LAMNOPPTDASE.
DR PROSITE: PS00631; CYTOSOL_AP; 1.
KW Hydrolyase; Aminopeptidase; Manganese; Complete proteome.
FT METAL 280 280 MANGANESE 2 (BY SIMILARITY).
FT METAL 285 285 MANGANESE 1 AND 2 (BY SIMILARITY).
FT METAL 303 303 MANGANESE 2 (BY SIMILARITY).
FT METAL 362 362 MANGANESE 1 (BY SIMILARITY).
FT METAL 364 364 MANGANESE 1 AND 2 (BY SIMILARITY).
FT ACT_SITE 292 292 POTENTIAL.
FT ACT_SITE 366 366 POTENTIAL.
SQ SEQUENCE 524 AA; 55609 MW; 0D6161A4D814C75D CRC64;

Query Match 89.7%; Score 26; DB 1; Length 524;
Best Local Similarity 80.0%; Pred. No. 4.8e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
Db 37 TRPR 41

Search completed: March 3, 2004, 12:17:40
Job time : 14 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: March 3, 2004, 12:15:23 ; Search time 20 Seconds
(without alignments)
24.048 Million cell updates/sec

Title: US-09-871-974-2

Perfect score: 29

Sequence: 1 TKPPR 5

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database :

PIR 78:*

1: PIR1:*

2: PIR2:*

3: PIR3:*

4: PIR4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	29	100.0	142	2 C82829	conserved hypothet
2	29	100.0	147	2 AH3157	conserved hypothet
3	29	100.0	191	2 T47369	hypothetical prote
4	29	100.0	201	2 S76026	hypothetical prote
5	29	100.0	222	2 G98129	hypothetical prote
6	29	100.0	226	2 T25360	hypothetical prote
7	29	100.0	278	2 S75883	hypothetical prote
8	29	100.0	283	2 T18299	hypothetical prote
9	29	100.0	350	2 AE3171	ATP-dependent DNA
10	29	100.0	351	2 T29369	hypothetical prote
11	29	100.0	370	2 AC0173	probable iron-sulf
12	29	100.0	375	2 D84715	probable protein k
13	29	100.0	410	2 S38238	hypothetical prote
14	29	100.0	415	2 F96499	hypothetical prote
15	29	100.0	415	2 C84829	hypothetical prote
16	29	100.0	442	2 S61165	hypothetical prote
17	29	100.0	493	2 S73752	hypothetical prote
18	29	100.0	535	2 T21775	hypothetical prote
19	29	100.0	604	2 S56027	hypothetical prote
20	29	100.0	615	2 JC7576	transcription fact
21	29	100.0	619	2 A43361	Sts-related transc
22	29	100.0	722	2 T22359	hypothetical prote
23	29	100.0	816	2 T45684	hypothetical prote
24	29	100.0	919	2 A41275	DNA ligase (ATP) (
25	29	100.0	941	2 T51135	ligand-gated chann
26	29	100.0	962	2 D86186	hypothetical prote
27	29	100.0	976	2 T51137	ionotropic glutama
28	29	100.0	1235	1 FWBYH	potassium transpor
29	29	100.0	1241	2 JU0466	potassium transpor

hypothetical prote
hypothetical prote
hypothetical prote
hypothetical prote
hypothetical prote
conserved hypothet
hypothetical prote
hypothetical prote
secretin precursor
vascular endotheli
ribosomal protein
hypothetical prote
hypothetical prote
hypothetical prote
transposase homolo
conserved homolo
hypothetical prote
hypothetical prote
acidic endoprotein
major core protein
hypothetical prote
guanylate kinase
guanylate kinase
guanylate kinase N
AcOrf-132 protein
ACMPV orf132 - Bo
hypothetical prote
hypothetical prote
tRNA pseudouridine
hypothetical prote
spermidine synthas
hypothetical prote
probable dihydrodi
hypothetical prote
putative transcrip
LuxA-related prote
calcium-binding pr
genome polyprotein
hypothetical prote
ATP-dependent DNA
site-specific DNA
type II DNA modifi
hypothetical prote
brefeldin A estera
tRNA (5-methylamin
probable GTP bindi
cholecystokinin ty
hypothetical prote
ATP-dependent RNA
Glycoprotein D pre
Glycoprotein D pre
55K erythrocyte me
homeotic protein H
probable cytosol a
phosphoinositide-s
oligopeptide trans
gag polyprotein -
hypothetical prote
phosphoinositide-s
phosphoinositide-s
phosphoinositide-s
phosphoinositide-s
conserved hypothet
phosphoinositide-s
phosphoinositide-s
phosphoinositide-s
phosphoinositide-s
phospholipase C1
phosphoinositide-s
phosphoinositide-s
phosphoinositide-s
phosphoinositide-s

submitted to GenBank, June 2000

A;Authors: Ferreira, V.C.A.; Ferro, J.A.; Fraga, J.S.; Franca, S.C.; Franco, M.C.; Froil
J.D.; Junqueira, M.L.; Kemper, E.L.; Kitajima, J.P.; Krieger, J.E.; Kuramae, E.E.; Lai
Chado, M.A.; Madeira, E.M.F.; Madeira, H.M.F.; Marino, C.L.; Marques, M.V.; Martins,
A;Authors: Martins, E.M.F.; Matsukuma, A.Y.; Menck, C.F.M.; Miracca, E.C.; Miyaki, C.Y
F.G.; Nunes, L.R.; Oliveira, M.A.; de Oliveira, M.C.; de Oliveira, R.C.; Palmieri, D
Rodrigues, V.; Rosa, A.J. de M.; de Rosa Jr., V.E.; de Sa, R.G.; Santelli, R.V.; Savas
A;Authors: da Silva, A.C.R.; da Silva, F.R.; da Silva, A.M.; Silva Jr., W.A.; da Silve
M.; Tshako, M.H.; Vallada, H.; Van Sluys, M.A.; Verjovski-Almeida, S.; Vettore, A.L.;

A;Reference number: A59328
A;Contents: annotation
C;Genetics:
A;Gene: XF0240

Query Match 100.0%; Score 29; DB 2; Length 142;
Best Local Similarity 100.0%; Pred. No. 64;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 87 TKPPR 91

RESULT 2
AH3157
conserved hypothetical protein Atu4884 [imported] - Agrobacterium tumefaciens (strain
C;Species: Agrobacterium tumefaciens
C;Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 18-Nov-2002
C;Accession: AH3157
R;Wood, D.W.; Sebail, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woo,
erage, G.; Gillet, W.; Grant, C.; Guenther, D.; Kutyavin, T.; Levy, R.; Li, M.; McCle
; Karp, P.; Romero, P.; Zhang, S.
Science 294, 2317-2323, 2001
A;Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamm
ster, E.W.
A;Title: The Genome of the Natural Genetic Engineer Agrobacterium tumefaciens C58.
A;Reference number: AB2577; MUID:21608550; PMID:11743193
A;Accession: AH3157
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-147 <KUR>
A;Cross-references: GB:AE008689; PIDN:AA45678.1; PID:gi7743404; GSPDB:GN00187
A;Experimental source: strain C58 (Dupont)
C;Genetics:
A;Gene: Atu4884
A;Map position: linear chromosome

Query Match 100.0%; Score 29; DB 2; Length 147;
Best Local Similarity 100.0%; Pred. No. 67;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 2 TKPPR 6

RESULT 3
T47369
hypothetical protein F7M19.120 - Arabidopsis thaliana
C;Species: Arabidopsis thaliana (mouse-ear cross)
C;Date: 20-Apr-2000 #sequence_revision 20-Apr-2000 #text_change 20-Apr-2000
C;Accession: T47369
R;Nyakatura, G.; Fartmann, B.; Dauner, D.; Sterr, W.; Holland, R.; Weichselgartner, M
Mayer, K.F.X.
submitted to the Protein Sequence Database, April 2000
A;Reference number: Z24458
A;Accession: T47369
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-191 <NYA>
A;Cross-references: EMBL:AL138643
A;Experimental source: cultivar Columbia; BAC clone F7M19
C;Genetics:

phospholipase C2 (

nuclear phosphoro

72K mitochondrial

hypothetical prote

hypothetical prote

ascites sialoglyco

hypothetical prote

probable formate d

exoribonuclease (i

exoribonuclease RN

hypothetical prote

HQRP1 protein - hu

probable cation tr

hypothetical prote

nitrate reductase

thrombospondin 4 p

hypothetical prote

translation elonga

dna exoribonuclease

hypothetical prote

eph receptor tyros

proline dehydrogen

protein F18014.19

hypothetical prote

immediate-early pr

immediate-early pr

ascites sialoglyco

voltage-dependent

voltage-dependent

calcium channel pr

calcium channel al

Munc13-3 protein -

calcium channel pr

genome polyprotein

ALR protein - huma

ALR protein - huma

microtubule-associ

cytochrome-c oxida

probable formylmet

ig lambda chain V-

hypothetical prote

hypothetical prote

probable outer mem

rubredoxin II - ps

zein, 27K - maize

gamma-coixin, 22K,

103 26 89.7 605 2 T50842

104 26 89.7 609 2 A43906

105 26 89.7 624 2 A36682

106 26 89.7 624 2 T21072

107 26 89.7 666 2 B70803

108 26 89.7 744 2 A43353

109 26 89.7 747 2 S69557

110 26 89.7 759 2 T35749

111 26 89.7 759 2 AG2736

112 26 89.7 784 2 AG7517

113 26 89.7 784 2 T33098

114 26 89.7 846 1 Q055C3

115 26 89.7 905 2 C07058

116 26 89.7 919 2 T29581

117 26 89.7 925 2 A36328

118 26 89.7 961 1 TSHUP4

119 26 89.7 965 2 T21073

120 26 89.7 975 2 T08606

121 26 89.7 983 2 T39902

122 26 89.7 991 2 S43891

123 26 89.7 1057 2 S09112

124 26 89.7 1122 2 T42400

125 26 89.7 1127 2 T28435

126 26 89.7 1158 2 E86327

127 26 89.7 1160 2 T00272

128 26 89.7 1160 2 A45344

129 26 89.7 1460 1 EDB2IF

130 26 89.7 1630 2 A53577

131 26 89.7 2139 2 A44467

132 26 89.7 2143 2 JH0427

133 26 89.7 2166 2 S11339

134 26 89.7 2171 2 S05054

135 26 89.7 2207 2 T42759

136 26 89.7 2220 2 A45290

137 26 89.7 3033 1 GNWVJ8

138 26 89.7 4957 2 T03455

139 26 89.7 5262 2 T03454

140 26 89.7 5327 2 T13564

141 25 86.2 14 2 S65392

142 25 86.2 69 1 B35537

143 25 86.2 105 2 E72599

144 25 86.2 111 1 L2HWN

145 25 86.2 126 2 H72614

146 25 86.2 133 2 E70690

147 25 86.2 168 2 C83505

148 25 86.2 173 1 RUPSP0

149 25 86.2 181 2 S22950

150 25 86.2 199 2 S23635

ALIGNMENTS

RESULT 1
C82829
conserved hypothetical protein XF0240 [imported] - Xylella fastidiosa (strain 9a5c)

C;Species: Xylella fastidiosa

C;Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 20-Aug-2000

C;Accession: C82829

R;anonymous, The Xylella fastidiosa Consortium of the Organization for Nucleotide Sequen

Nature 406, 151-157, 2000

A;Title: The genome sequence of the plant pathogen Xylella fastidiosa.

A;Reference number: AB2515; MUID:20365717; PMID:10910347

A;Note: for a complete list of authors see reference number A59328 below

A;Accession: C82829

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-142 <SIM>

A;Cross-references: GB:AE003878; GB:AE003849; NID:G9105052; PIDN:AAF83053.1; GSPDB:GN001

A;Experimental source: strain 9a5c

R;Simpson, A.J.G.; Reinach, F.C.; Arruda, P.; Abreu, F.A.; Acencio, M.; Alvarenga, R.; A

Briones, M.R.S.; Bueno, M.R.P.; Camargo, A.A.; Camargo, L.E.A.; Carraro, D.M.; Carreir, H

as-Neto, E.; Docena, C.; El-Dorry, H.; Facincani, A.P.; Ferreira, A.J.S.

A;Map position: 3
A;Note: F7M19.120

Query Match 100.0%; Score 29; DB 2; Length 191;
Best Local Similarity 100.0%; Pred. No. 87;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|
|
|
|
|
Db 78 TKPPR 82

RESULT 4

S76026
hypothetical protein - Synecocystis sp. (strain PCC 6803)
C;Species: Synecocystis sp.
A;Variety: PCC 6803
C;Date: 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change 20-Jun-2000
C;Accession: S76026
R;Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.; Nakamura, Y.; Miyajima, N.; O. K.; Okumura, S.; Shimpo, S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.; Yasuda
DNA Res. 3, 109-136, 1996
A;Title: Sequence analysis of the genome of the unicellular cyanobacterium Synecocystis
s.
A;Reference number: S74322; MUID:97061201; PMID:8905231
A;Accession: S76026
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-201 <KAN>
A;Cross-references: EMBL:D64006; GB:AB001339; NID:g1001291; PIDN:BAI0873.1; PID:g100138
A;Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996
C;Superfamily: conserved hypothetical protein YDR196c

Query Match 100.0%; Score 29; DB 2; Length 201;
Best Local Similarity 100.0%; Pred. No. 91;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|
|
|
|
|
Db 3 TKPPR 7

RESULT 5

G98129
hypothetical protein AGR_L19 [imported] - Agrobacterium tumefaciens (strain C58, Cereon
C;Species: Agrobacterium tumefaciens
C;Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 18-Nov-2002
C;Accession: G98129
R;Goodner, B.; Hinkle, G.; Gattung, S.; Miller, N.; Blanchard, M.; Qurollo, B.; Goldman,
A.; Liu, F.; Wollam, C.; Allinger, M.; Doughty, D.; Scott, C.; Lappas, C.; Markelz, B.;
Science 294, 2323-2328, 2001
A;Title: Genome Sequence of the Plant Pathogen and Biotechnology Agent Agrobacterium tum
A;Reference number: A97359; MUID:21608551; PMID:11743194
A;Accession: G98129
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-222 <KUR>
A;Cross-references: GB:AF007870; PIDN:AAK88561.1; PID:g15158270; GSPDB:GN00170
C;Genetics:
A;Gene: AGR_L19
A;Map position: linear chromosome

Query Match 100.0%; Score 29; DB 2; Length 222;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|
|
|
|
|
Db 77 TKPPR 81

RESULT 6

T25360

hypothetical protein T27E7.1 - Caenorhabditis elegans
C;Species: Caenorhabditis elegans
C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
C;Accession: T25360
R;Cummings, P.
submitted to the EMBL Data Library, November 1996
A;Reference number: Z20023
A;Accession: T25360
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 1-226 <WIL>
A;Cross-references: EMBL:Z82284; PIDN:CAB05287.1; GSPDB:GN00022; CESP:T27E7.1
A;Experimental source: Clone T27E7
C;Genetics:
A;Gene: CESP:T27E7.1
A;Map position: 4
A;Introns: 20/1; 43/1; 101/3; 169/3

Query Match 100.0%; Score 29; DB 2; Length 226;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|
|
|
|
|
Db 75 TKPPR 79

RESULT 7

S75883
hypothetical protein slr1169 - Synecocystis sp. (strain PCC 6803)
C;Species: Synecocystis sp.
A;Variety: PCC 6803
C;Date: 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change 08-Oct-1999
C;Accession: S75883
R;Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.; Nakamura, Y.; Miyajima, N.; O. K.; Okumura, S.; Shimpo, S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.; Yasu
DNA Res. 3, 109-136, 1996
A;Title: Sequence analysis of the genome of the unicellular cyanobacterium Synecocyst
s.
A;Reference number: S74322; MUID:97061201; PMID:8905231
A;Accession: S75883
A;Status: nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-278 <KAN>
A;Cross-references: EMBL:P90913; GB:AB001339; NID:g1653348; PIDN:BAI18342.1; PID:dl019
A;Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996

Query Match 100.0%; Score 29; DB 2; Length 278;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|
|
|
|
|
Db 240 TKPPR 244

RESULT 8

T18299
hypothetical protein - Entamoeba histolytica (fragment)
C;Species: Entamoeba histolytica
C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 21-Jul-2000
C;Accession: T18299
R;Evans, J.D.
Proc. Natl. Acad. Sci. U.S.A. 92, 6518-6521, 1995
A;Title: Relatedness threshold for the production of female sexuals in colonies of a p
A;Reference number: Z18867; MUID:95327678; PMID:7604025
A;Accession: T18299
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 1-283 <EVA>
A;Cross-references: EMBL:L39933; NID:g6478875; PIDN:AAC41578.1; PID:g675517

Query Match 100.0%; Score 29; DB 2; Length 283;

```

Best Local Similarity 100.0%; Pred. No. 1.3e+02; Indels 0; Gaps 0;
Matches 5; Conservative 0; Mismatches 0;

QY 1 TKPPR 5
    |||||
Db 208 TKPPR 212

RESULT 9
AE3171
ATP-dependent DNA ligase Atu5097 [imported] - Agrobacterium tumefaciens (strain C58, Dup
C:Species: Agrobacterium tumefaciens
C>Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 18-Nov-2002
C:Accession: AE3171
R:Wood, D.W.; Setubal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woc, I
erage, G.; Gillet, W.; Grant, C.; Guenther, D.; Kutyavin, T.; Levy, R.; Li, M.; McClell
; Karp, P.; Romero, P.; Zhang, S.
Science 294, 2317-2323, 2001
A:Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-kamm,
ster, E.W.
A:Title: The Genome of the Natural Genetic Engineer Agrobacterium tumefaciens C58.
A:Reference number: AB2577; MUID:21608550; PMID:11743193
A:Accession: AE3171
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-350 <KUR>
A:Cross-references: GB:AB008687; PIDN:AAL45787.1; PID:gl7743523; GSPDB:GN00188
A:Experimental source: strain C58 (Dupont)
C:Genetics:
A:Gene: Atu5097
A:Genome: plasmid

Query Match 100.0%; Score 29; DB 2; Length 350;
Best Local Similarity 100.0%; Pred. No. 1.6e+02; Indels 0; Gaps 0;
Matches 5; Conservative 0; Mismatches 0;

QY 1 TKPPR 5
    |||||
Db 2 TKPPR 6

RESULT 10
T29369
Hypothetical protein ZC404.8 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
C:Accession: T29369
R:Bentley, D.; Lee, T.T.
submitted to the EMBL Data Library, April 1996
A:Description: The sequence of C. elegans cosmid ZC404.
A:Reference number: Z20614
A:Accession: T29369
A:Status: preliminary; translated from GB/EMBL/DD5J
A:Molecule type: DNA
A:Residues: 1-351 <BEN>
A:Cross-references: EMBL:U55363; PIDN:AAA97963.1; GSPDB:GN00023; CESP:ZC404.8
A:Experimental source: strain Bristol N2; clone ZC404
C:Genetics:
A:Gene: CESP:ZC404.8
A:Map position: 5
A:Introns: 17/2; 52/2; 73/2; 312/1

Query Match 100.0%; Score 29; DB 2; Length 351;
Best Local Similarity 100.0%; Pred. No. 1.6e+02; Indels 0; Gaps 0;
Matches 5; Conservative 0; Mismatches 0;

QY 1 TKPPR 5
    |||||
Db 199 TKPPR 203

RESULT 11
AC0173
probable iron-sulfur binding protein YP01417 [imported] - Yersinia pestis (strain CO92
C:Species: Yersinia pestis
C>Date: 02-Nov-2001 #sequence_revision 02-Nov-2001 #text_change 02-Nov-2001
C:Accession: AC0173
R:Parkhill, J.; Wren, B.W.; Thomson, N.R.; Titball, R.W.; Holden, M.T.G.; Prentice, M.
deno-Tarraga, A.M.; Chillingworth, T.; Cronin, A.; Davies, R.M.; Davis, P.; Dougan, G.
il, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.; Whitehead, S.; Barrell
Nature 413, 523-527, 2001
A:Title: Genome sequence of Yersinia pestis, the causative agent of plague.
A:Reference number: AB0001; MUID:21470413; PMID:11586360
A:Accession: AC0173
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-370 <KUR>
A:Cross-references: GB:AL590842; PIDN:CAC90246.1; PID:gl5979466; GSPDB:GN00175
C:Genetics:
A:Gene: YP01417

Query Match 100.0%; Score 29; DB 2; Length 370;
Best Local Similarity 100.0%; Pred. No. 1.7e+02; Indels 0; Gaps 0;
Matches 5; Conservative 0; Mismatches 0;

QY 1 TKPPR 5
    |||||
Db 266 TKPPR 270

RESULT 12
D84715
Probable protein kinase [imported] - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C>Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
C:Accession: D84715
R:Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.
M.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; VanAken, S.E.; Umayam, L.; Tallon,
eues, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter,
Nature 402, 761-768, 1999
A:Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.
A:Reference number: A84420; MUID:20083487; PMID:10617157
A:Accession: D84715
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-375 <STO>
A:Cross-references: GB:AE002093; NID:g3201626; PIDN:AAC20735.1; GSPDB:GN00139
C:Genetics:
A:Gene: At2g31010
A:Map position: 2

Query Match 100.0%; Score 29; DB 2; Length 375;
Best Local Similarity 100.0%; Pred. No. 1.7e+02; Indels 0; Gaps 0;
Matches 5; Conservative 0; Mismatches 0;

QY 1 TKPPR 5
    |||||
Db 121 TKPPR 125

RESULT 13
S38238
Hypothetical protein - Coxiella burnetii
C:Species: Coxiella burnetii
C>Date: 20-Feb-1995 #sequence_revision 20-Feb-1995 #text_change 08-Oct-1999
C:Accession: S38238
R:Thiele, D.; Willems, H.; Haas, M.; Krauss, H.
submitted to the EMBL Data Library, October 1993
A:Reference number: S38215
A:Accession: S38238
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-410 <THI>
A:Cross-references: EMBL:X75356; NID:ig407370; PIDN:CAA53126.1; PID:g407394

Query Match 100.0%; Score 29; DB 2; Length 410;

```

```

Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 102 TKPPR 106

RESULT 14
F96499
hypothetical protein T10P12.9 [imported] - Arabidopsis thaliana
C;Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001
R;Accession: F96499
R;Theologos, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Alonso,
Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Conway, A.R.; Creasy, T.H.; Dewar, K.;
ansen, N.F.; Hughes, B.; Huizar, L.
Nature 408, 816-820, 2000
A;Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Kim, C.;
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Marziali,
Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.;
A;Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallon,
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A;Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
A;Reference number: A86141; MUID:21016719; PMID:11130712
A;Accession: F96499
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-415 <STO>
A;Cross-references: GB:AE005173; NID:G5080765; PIDN:AA039275.1; GSPDB:GN00141
C;Genetics:
A;Gene: T10P12.9
A;Map position: 1

Query Match 100.0%; Score 29; DB 2; Length 415;
Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 309 TKPPR 313

RESULT 15
C84829
hypothetical protein At2g40420 [imported] - Arabidopsis thaliana
C;Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
C;Accession: C84829
R;Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.;
M.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; VanAken, S.E.; Umayam, L.; Tallon, L.;
euss, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter, J.
Nature 402, 761-768, 1999
A;Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.
A;Reference number: A84420; MUID:20083487; PMID:10617197
A;Accession: C84829
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-415 <STO>
A;Cross-references: GB:AE002093; NID:G6598346; PIDN:AAB87575.2; GSPDB:GN00139
C;Genetics:
A;Gene: At2g40420
A;Map position: 2

Query Match 100.0%; Score 29; DB 2; Length 415;
Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 169 TKPPR 173

RESULT 16
S61165
hypothetical protein YDR370c - yeast (Saccharomyces cerevisiae)
N;Alternate names: hypothetical protein D9481.14
C;Species: Saccharomyces cerevisiae
C;Date: 23-Feb-1996 #sequence_revision 01-Mar-1996 #text_change 19-Apr-2002
C;Accession: S61165
R;Ding, H.
submitted to the EMBL Data Library, June 1995
A;Description: The sequence of S. cerevisiae cosmid 9481.
A;Reference number: S61159
A;Accession: S61165
A;Molecule type: DNA
A;Residues: 1-442 <DIN>
A;Cross-references: EMBL:U29373; NID:G649184; PID:G849191; GSPDB:GN00004; MIPS:YDR370c
A;Experimental source: strain S288C (AB972)
C;Genetics:
A;Gene: MIPS:YDR370c
A;Cross-references: SGD:S0002778
A;Map position: 4R
C;Superfamily: Saccharomyces cerevisiae hypothetical protein YDR370c

Query Match 100.0%; Score 29; DB 2; Length 442;
Best Local Similarity 100.0%; Pred. No. 2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 81 TKPPR 85

RESULT 17
S73752
hypothetical protein A05_orf493 - Mycoplasma pneumoniae (strain ATCC 29342)
C;Species: Mycoplasma pneumoniae
A;Variety: ATCC 29342
C;Date: 27-Feb-1997 #sequence_revision 25-Apr-1997 #text_change 15-Sep-2000
C;Accession: S73752
R;Himmelreich, R.; Hilbert, H.; Plagens, H.; Pirkl, E.; Li, B.C.; Herrmann, R.
Nucleic Acids Res. 24, 4420-4449, 1996
A;Title: Complete sequence analysis of the genome of the bacterium Mycoplasma pneumoniae
A;Reference number: S73327; MUID:97105885; PMID:8948633
A;Accession: S73752
A;Status: preliminary; nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-493 <HIM>
A;Cross-references: EMBL:AE000041; GB:U00089; NID:G1674104; PIDN:AAB96074.1; PID:G16741
A;Note: the nucleotide sequence was submitted to the EMBL Data Library, November 1996
C;Genetics:
A;Genetic code: SGC3
C;Superfamily: Mycoplasma pneumoniae hypothetical protein H08_orf445

Query Match 100.0%; Score 29; DB 2; Length 493;
Best Local Similarity 100.0%; Pred. No. 2.2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 462 TKPPR 466

RESULT 18
T21775
hypothetical protein F35E2.5 - Caenorhabditis elegans
C;Species: Caenorhabditis elegans
C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
C;Accession: T21775
R;Lennard, N.
submitted to the EMBL Data Library, November 1996
A;Reference number: Z19471
A;Accession: T21775
A;Status: preliminary; translated from GB/EMBL/DBDJ
A;Molecule type: DNA

```


A;Residues: 1-575 <WIL>
 A;Cross-references: EMBL:281528; PIDN:CA804281.1; GSPDB:GN00019; CESP:F35E2.5
 A;Experimental source: clone F35E2
 C;Genetics:
 A;Gene: CESP:F35E2.5
 A;Map position: 1
 A;Introns: 44/1; 109/2; 209/3; 250/1; 367/1; 413/1; 480/3; 508/3

Query Match 100.0%; Score 29; DB 2; Length 575;
 Best Local Similarity 100.0%; Pred. No. 2.6e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
 |||||
 Db 284 TKPPR 288

RESULT 19

S56027
 hypothetical protein YUL083w - yeast (Saccharomyces cerevisiae)
 N;Alternate names: hypothetical protein YJ002
 C;Species: Saccharomyces cerevisiae
 C;Date: 05-May-1995 #sequence_revision 08-Sep-1995 #text_change 19-Apr-2002
 C;Accession: S56027; S56850; S57742
 R;Mitsuga, T.; Schaaff-Gerstenschlaeger, I.; Chaiwatizis, N.; Baur, A.; Boles, E.; Fournie
 Yeast 11, 681-689, 1995
 A;Title: Sequence analysis of a 33.1 kb fragment from the left arm of Saccharomyces cere
 ter domain and a putative alpha-2-SCB-alpha-2 binding site.
 A;Reference number: S56016; MUID:96093911; PMID:7483841
 A;Accession: S56027
 A;Status: nucleic acid sequence not shown; translation not shown
 A;Molecule type: DNA
 A;Residues: 1-604 <MIO>
 A;Cross-references: EMBL:X83502; NID:G929861; PIDN:CA858487.1; PID:G929873
 A;Note: the nucleotide sequence was submitted to the EMBL Data Library, December 1994
 R;Mitsuga, T.; Schaaff-Gerstenschlaeger, I.; Baur, A.; Boles, E.; Chaiwatizis, N.; Fournie
 submitted to the Protein Sequence database, September 1995
 A;Reference number: S56855
 A;Accession: S56850
 A;Molecule type: DNA
 A;Residues: 1-604 <MIW>
 A;Cross-references: EMBL:249358; NID:G1008247; PIDN:CAA89376.1; PID:G1008248; MIPS:YJL08
 R;Sor, F.J.
 submitted to the EMBL Data Library, June 1995
 A;Reference number: S57731
 A;Accession: S57742
 A;Molecule type: DNA
 A;Residues: 1-604 <SOR>
 A;Cross-references: EMBL:X88851; NID:G995892; PIDN:CAA61318.1; PID:G995904
 C;Genetics:
 A;Cross-references: SGD:S0003619
 A;Map position: 10L

Query Match 100.0%; Score 29; DB 2; Length 604;
 Best Local Similarity 100.0%; Pred. No. 2.7e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
 |||||
 Db 151 TKPPR 155

RESULT 20

JC7576
 transcription factor Elf-1, type 1 - rat
 C;Species: Rattus norvegicus (Norway rat)
 C;Date: 30-Jun-2001 #sequence_revision 30-Jun-2001 #text_change 30-Jun-2001
 C;Accession: JC7576
 R;Nishiyama, C.; Takahashi, K.; Nishiyama, M.; Okumura, K.; Ra, C.; Ohtake, Y.; Yokota,
 Biosci. Biotechnol. Biochem. 64, 2601-2607, 2000
 A;Title: Splice isoforms of transcription factor Elf-1 affecting its regulatory function
 A;Reference number: JC7576; MUID: 21077473; PMID:11210123
 A;Contents: Mast cell line, RBL-2H3

A;Accession: JC7576
 A;Molecule type: mRNA
 A;Residues: 1-615 <NIS>
 A;Cross-references: DDBJ:AB030215
 C;Comment: This protein, as a key transcription factor for immune-related genes, has tr
 C;Genetics:
 A;Gene: Elf-1

Query Match 100.0%; Score 29; DB 2; Length 615;
 Best Local Similarity 100.0%; Pred. No. 2.8e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
 |||||
 Db 179 TKPPR 183

RESULT 21

A43361
 Ets-related transcription factor Elf-1 - human
 N;Alternate names: E74-like factor Elf-1
 C;Species: Homo sapiens (man)
 C;Date: 11-Feb-1993 #sequence_revision 11-Feb-1993 #text_change 07-May-1999
 C;Accession: A43361; A42122
 R;Laiden, J.M.; Wang, C.Y.; Petryniak, B.; Markovitz, D.M.; Nabel, G.J.; Thompson, C.B
 J. Virol. 66, 5890-5897, 1992
 A;Title: A novel Ets-related transcription factor, Elf-1, binds to human immunodeficien
 A;Reference number: A43361; MUID:92407982; PMID:1527846
 A;Accession: A43361
 A;Molecule type: mRNA
 A;Residues: 1-619 <LEI>
 A;Cross-references: GB:M82882
 R;Thompson, C.B.; Wang, C.Y.; Ho, I.C.; Bohjanen, P.R.; Petryniak, B.; June, C.H.; Mie
 Mol. Cell. Biol. 12, 1043-1053, 1992
 A;Title: cis-acting sequences required for inducible interleukin-2 enhancer function b
 A;Reference number: A42122; MUID:92186836; PMID:1545787
 A;Accession: A42122
 A;Status: preliminary; not compared with conceptual translation
 A;Molecule type: mRNA
 A;Residues: 204-282, 'G', 284-289 <THO>
 A;Experimental source: T-cells
 A;Note: sequence extracted from NCBI backbone (NCBI:P:88288)
 C;Genetics:
 A;Gene: GDB:ELF1
 A;Cross-references: GDB:131648
 A;Map position: lp36-lp36
 C;Superfamily: ets DNA-binding domain homology
 C;Keywords: DNA binding; transcription regulation
 F;210-290/Domain: ets DNA-binding domain homology <ETS>

Query Match 100.0%; Score 29; DB 2; Length 619;
 Best Local Similarity 100.0%; Pred. No. 2.8e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
 |||||
 Db 180 TKPPR 184

RESULT 22

T22359
 hypothetical protein F47G4.4 - Caenorhabditis elegans
 C;Species: Caenorhabditis elegans
 C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
 C;Accession: T22359
 R;White, S.
 submitted to the EMBL Data Library, September 1997
 A;Reference number: Z19553
 A;Accession: T22359
 A;Status: preliminary; translated from GB/EMBL/DBJ
 A;Molecule type: DNA
 A;Residues: 1-722 <WIL>
 A;Cross-references: EMBL:Z99171; PIDN:CA816313.1; GSPDB:GN00019; CESP:F47G4.4

A;Experimental source: clone F47G4

C;Genetics:

A;Gene: CESP.F47G4.4

A;Map position: 1

A;Introns: 5/1; 43/3; 129/3; 176/3; 226/1; 259/3; 319/3; 365/3; 465/2; 576/1; 631/1; 688

Query Match 100.0%; Score 29; DB 2; Length 722;

Best Local Similarity 100.0%; Pred. No. 3.3e+02;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5

DB 402 TKPPR 406

RESULT 23

T45684

hypothetical protein F14P22.230 - Arabidopsis thaliana

C;Species: Arabidopsis thaliana (mouse-ear cress)

C;Date: 04-Feb-2000 #sequence_revision 04-Feb-2000 #text_change 04-Mar-2000

C;Accession: T45684

R;D'Angelo, M.; Vezzi, A.; Modesto, D.; Pigazzi, M.; Valle, G.; Mewes, H.W.; Lemcke, K.

submitted to the Protein Sequence Database, January 2000

A;Reference number: Z23011

A;Accession: T45684

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-816 <DAN>

A;Cross-references: EMBL:AL137082

A;Experimental source: cultivar Columbia; BAC clone F14P22

C;Genetics:

A;Map position: 3

A;Introns: 83/3; 151/3; 209/3; 296/2; 336/2; 366/2; 398/2; 429/2; 469/3; 562/1; 603/2; 6

A;Note: F14P22.230

C;Superfamily: unassigned Ser/Thr or Tyr-specific protein kinases; protein kinase homolo

Query Match 100.0%; Score 29; DB 2; Length 816;

Best Local Similarity 100.0%; Pred. No. 3.7e+02;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5

DB 625 TKPPR 629

RESULT 24

A41275

DNA ligase (ATP) (EC 6.5.1.1) I - human

N;Alternate names: DNA joinase; DNA repair enzyme; polydeoxyribonucleotide synthase (ATP

C;Species: Homo sapiens (man)

C;Date: 16-Sep-1992 #sequence_revision 16-Sep-1992 #text_change 03-Feb-2003

C;Accession: A36048; A41275

R;Barnes, D.E.; Johnston, L.H.; Kodama, K.; Tomkinson, A.E.; Lasko, D.D.; Lindahl, T.

Proc. Natl. Acad. Sci. U.S.A. 87, 6679-6683, 1990

A;Title: Human DNA ligase I cDNA: cloning and functional expression in Saccharomyces cer

A;Reference number: A36048; MUID:90370849; PMID:2204063

A;Accession: A36048

A;Molecule type: mRNA

A;Residues: 1-919 <BAR>

A;Cross-references: GB:M36067; NID:g187142; PIDN:AAA59518.1; PID:g187143

R;Petrini, J.H.J.; Huwiler, K.G.; Weaver, D.T.

Proc. Natl. Acad. Sci. U.S.A. 88, 7615-7619, 1991

A;Title: A wild-type DNA ligase I gene is expressed in Bloom's syndrome cells.

A;Reference number: A41275; MUID:91352039; PMID:1881902

A;Accession: A41275

A;Molecule type: mRNA

A;Residues: 716-753 <PEP>

C;Genetics:

A;Gene: GDB:LIG1

A;Cross-references: GDB:127274; OMIM:126391

A;Map position: 19q13.3-19q13.3

C;Keywords: DNA repair; ligase; phosphoprotein

F;568/Active site: Lys (covalent AMP-binding) #status predicted

Query Match 100.0%; Score 29; DB 2; Length 919;

Best Local Similarity 100.0%; Pred. No. 4.2e+02;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5

DB 218 TKPPR 222

RESULT 25

T51135

ligand-gated channel-like protein precursor [imported] - Arabidopsis thaliana

C;Species: Arabidopsis thaliana (mouse-ear cress)

C;Date: 28-Jul-2000 #sequence_revision 28-Jul-2000 #text_change 28-Jul-2000

C;Accession: T51135

R;Meyerhoff, O.; Hedrich, R.; Becker, D.

submitted to the EMBL Data Library, July 1999

A;Description: Characterization of ligand-gated channel-like proteins in higher plants

A;Reference number: Z25308

A;Accession: T51135

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: mRNA

A;Residues: 1-941 <MEY>

A;Cross-references: EMBL:AF167355; PIDN:AAD47833.1

A;Experimental source: cultivar Columbia

C;Genetics:

A;Gene: GLUR3

A;Map position: 1

Query Match 100.0%; Score 29; DB 2; Length 941;

Best Local Similarity 100.0%; Pred. No. 4.3e+02;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5

DB 468 TKPPR 472

RESULT 26

D86186

hypothetical protein [imported] - Arabidopsis thaliana

C;Species: Arabidopsis thaliana (mouse-ear cress)

C;Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001

C;Accession: D86186

R;Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Alonso

Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Conway, A.R.; Creasy, T.H.; Dewar, K.

ansen, N.P.; Hughes, B.; Huizar, L.

Nature 408, 816-820, 2000

A;Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, B.; Kim, C.

C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Marziani

Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.

A;Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallon,

ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.

A;Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.

A;Reference number: A86141; MUID:21016719; PMID:11130712

A;Accession: D86186

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-962 <STO>

A;Cross-references: GB:AE005172; NID:g2388577; PIDN:AAB71458.1; GSPDB:GN00141

C;Genetics:

A;Map position: 1

Query Match 100.0%; Score 29; DB 2; Length 962;

Best Local Similarity 100.0%; Pred. No. 4.4e+02;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5

DB 468 TKPPR 472

RESULT 27

IS1137
ionotropic glutamate receptor homolog GLR4 [imported] - Arabidopsis thaliana
C/Species: Arabidopsis thaliana (mouse-ear cress)
C/Date: 28-Jul-2000 #sequence_revision 28-Jul-2000 #text_change 28-Jul-2000
C/Accession: T51137
R/Davenport, R.J.; Kiegle, E.A.; Tester, M.
submitted to the EMBL Data Library, September 1999
A/Description: Cloning of an ionotropic glutamate receptor homolog from Arabidopsis thaliana
A/Reference number: Z5310
A/Accession: T51137
A/Status: preliminary; translated from GB/EMBL/DDRJ
A/Molecule type: mRNA
A/Residues: 1-976 <DAV>
A/Cross-references: EMBL:AF183932; PIDN:AAF01294.1
C/Genetics:
A/Gene: GLR4

Query Match 100.0%; Score 29; DB 2; Length 976;
Best Local Similarity 100.0%; Pred. No. 4.4e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 468 TKPPR 472

RESULT 28

FWYH
potassium transport protein TRK1, high-affinity - yeast (Saccharomyces cerevisiae)
N/Alternate names: protein J0693; protein YJL129c
C/Species: Saccharomyces cerevisiae
C/Date: 31-Mar-1991 #sequence_revision 31-Mar-1991 #text_change 27-Oct-2003
C/Accession: S05849; S56910
R/Gaber, R.F.; Styles, C.A.; Fink, G.R.
Mol. Cell. Biol. 8, 2848-2859, 1988
A/Title: TRK1 encodes a plasma membrane protein required for high-affinity potassium transport
A/Reference number: S05849; MUID:88302204; PMID:3043197
A/Accession: S05849
A/Molecule type: DNA
A/Residues: 1-1235 <GAB>
A/Cross-references: EMBL:M21328; NID:gl71803; PIDN:AAA34728.1; PID:gl71804
R/Czipluch, C.; Kordes, E.; Pujol, A.; Jauniaux, J.C.
submitted to the Protein Sequence Database, September 1995
A/Reference number: S56891
A/Accession: S56910
A/Molecule type: DNA
A/Residues: 1-1235 <CZI>
A/Cross-references: EMBL:Z49404; NID:gl008329; PIDN:CAA89424.1; PID:gl008330; GSPDB:GN00
C/Genetics:
A/Gene: SGD:TRK1; MIPS:YJL129c
A/Cross-references: SGD:S0003665; MIPS:YJL129c
C/Superfamily: potassium transport protein TRK1/TRK2
C/Keywords: ion transport; potassium transport; transmembrane protein
F/50-70/Domain: transmembrane #status predicted <TM01>
F/72-96/Domain: transmembrane #status predicted <TM02>
F/107-127/Domain: transmembrane #status predicted <TM03>
F/178-806/Domain: transmembrane #status predicted <TM04>
F/819-840/Domain: transmembrane #status predicted <TM05>
F/844-864/Domain: transmembrane #status predicted <TM06>
F/868-889/Domain: transmembrane #status predicted <TM07>
F/904-924/Domain: transmembrane #status predicted <TM08>
F/929-949/Domain: transmembrane #status predicted <TM09>
F/1084-1104/Domain: transmembrane #status predicted <TM10>
F/1117-1137/Domain: transmembrane #status predicted <TM11>

Query Match 100.0%; Score 29; DB 1; Length 1235;
Best Local Similarity 100.0%; Pred. No. 5.6e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 438 TKPPR 442

RESULT 29

JU0466
potassium transport protein TRK1, high-affinity - yeast (Saccharomyces cerevisiae) (str N/Alternate names: protein YJL129c
C/Species: Saccharomyces cerevisiae
A/Variety: Saccharomyces uvarum
C/Date: 13-Jul-1990 #sequence_revision 28-Oct-1994 #text_change 27-Oct-2003
C/Accession: JU0466
R/Anderson, J.A.; Best, L.A.; Gaber, R.F.
Gene 99, 39-46, 1991
A/Title: Structural and functional conservation between the high-affinity K⁺ transporter
A/Reference number: JU0466; MUID:91216443; PMID:2022322
A/Accession: JU0466
A/Molecule type: DNA
A/Residues: 1-1241 <AND>
A/Cross-references: GB:M57508; NID:gl71640; PIDN:AAA34661.1; PID:gl71641
A/Note: the source is designated as Saccharomyces uvarum
C/Genetics:
A/Gene: SGD:TRK1
A/Cross-references: SGD:S0003665; MIPS:YJL129c
A/Map position: 10L
C/Superfamily: potassium transport protein TRK1/TRK2
C/Keywords: ion transport; potassium transport; transmembrane protein
F/50-70/Domain: transmembrane #status predicted <TM01>
F/78-98/Domain: transmembrane #status predicted <TM02>
F/107-117/Domain: transmembrane #status predicted <TM03>
F/178-806/Domain: transmembrane #status predicted <TM04>
F/819-840/Domain: transmembrane #status predicted <TM05>
F/844-864/Domain: transmembrane #status predicted <TM06>
F/868-889/Domain: transmembrane #status predicted <TM07>
F/904-924/Domain: transmembrane #status predicted <TM08>
F/929-949/Domain: transmembrane #status predicted <TM09>
F/1084-1104/Domain: transmembrane #status predicted <TM10>
F/1117-1137/Domain: transmembrane #status predicted <TM11>

Query Match 100.0%; Score 29; DB 2; Length 1241;
Best Local Similarity 100.0%; Pred. No. 5.6e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 444 TKPPR 448

RESULT 30

G81008
hypothetical protein NMB2072 [imported] - Neisseria meningitidis (strain MC58 serogroup C/Species: Neisseria meningitidis
C/Date: 31-Mar-2000 #sequence_revision 31-Mar-2000 #text_change 19-Jan-2001
C/Accession: G81008
R/Tettelin, H.; Saunders, N.J.; Heidelberg, J.; Jeffries, A.C.; Nelson, K.E.; Eisen, J. Hickey, E.K.; Haft, D.H.; Salzberg, S.L.; White, O.; Fleischmann, R.D.; Dougherty, B.A. ri, H.; Qin, H.; Vamathevan, J.; Gill, J.; Scarlato, V.; Masignani, V.; Pizza, M.
Science 287, 1809-1815, 2000
A/Authors: Grandi, G.; Sun, L.; Smith, H.O.; Fraser, C.M.; Moxon, E.R.; Rappuoli, R.; A/Title: Complete genome sequence of Neisseria meningitidis serogroup B strain MC58.
A/Reference number: A81000; MUID:20175755; PMID:10710307
A/Accession: G81008
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-47 <TET>
A/Cross-references: GB:AB002557; GB:AB002098; NID:gl7227332; PIDN:AAF42391.1; PID:gl7227
A/Experimental source: serogroup B, strain MC58
C/Genetics:
A/Gene: NMB2072

Query Match 89.7%; Score 26; DB 2; Length 47;
Best Local Similarity 80.0%; Pred. No. 85;

Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|
Db 20 TRPPR 24

RESULT 31
AH2135
hypothetical protein asl2639 [imported] - Nostoc sp. (strain PCC 7120)
C/Species: Nostoc sp. PCC 7120
A/Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120
C/Date: 14-Dec-2001 #sequence_revision 14-Dec-2001 #text_change 09-Dec-2002
C/Accession: AH2135
R;Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasamoto, S.; Watanabe, A.; Iriiguchi, N.; Shimpo, S.; Sugimoto, M.; Takazawa, M.; Yamada, M.; Yasuda, M.; Tabata, S.
DNA Res. 8, 205-213, 2001
A/Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium Anabaena
A/Reference number: AB1807; MUID:21595285; PMID:11759840
A/Accession: AH2135
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-50 <KUR>
A/Cross-references: GB:BA000019; PIDN:BA074338.1; PID:g17131732; GSPDB:GN00179
A/Experimental source: strain PCC 7120
C/Genetics:
A/Gene: asl2639

Query Match 89.7%; Score 26; DB 2; Length 50;
Best Local Similarity 80.0%; Pred. No. 91;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|
Db 20 TKPPK 24

RESULT 32
C81143
hypothetical protein NMB0921 [imported] - Neisseria meningitidis (strain MC58 serogroup C)
C/Species: Neisseria meningitidis
C/Date: 31-Mar-2000 #sequence_revision 31-Mar-2000 #text_change 19-Jan-2001
C/Accession: C81143
R;Tettelin, H.; Saunders, N.J.; Heidelberg, J.; Jeffries, A.C.; Nelson, K.E.; Eisen, J.A.; Hickey, E.K.; Haft, D.H.; Salzberg, S.L.; White, O.; Fleischmann, R.D.; Dougherty, B.A.; Ri, H.; Qin, H.; Vamathevan, J.; Gill, J.; Scarlato, V.; Masignani, V.; Pizza, M.
Science 287, 1809-1815, 2000
A/Authors: Grandi, G.; Sun, L.; Smith, H.O.; Fraser, C.M.; Moxon, E.R.; Rappuoli, R.; Venter
A/Title: Complete genome sequence of Neisseria meningitidis serogroup B strain MC58.
A/Reference number: AB1000; MUID:20175755; PMID:10710307
A/Accession: C81143
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-66 <TET>
A/Cross-references: GB:AE002443; GB:AE002098; NID:g7226149; PIDN:AAF41329.1; PID:g722615
A/Experimental source: serogroup B, strain MC58
C/Genetics:
A/Gene: NMB0921

Query Match 89.7%; Score 26; DB 2; Length 66;
Best Local Similarity 80.0%; Pred. No. 1.2e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|
Db 39 TKPPK 43

RESULT 33
T36774
hypothetical protein SC128.04c - Streptomyces coelicolor
C/Species: Streptomyces coelicolor
C/Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Dec-1999

C/Accession: T36774
R;Seeger, K.; Harris, D.; Bentley, S.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.
Submitted to the EMBL Data Library, July 1999
A/Reference number: Z21574
A/Accession: T36774
A/Status: preliminary; translated from GB/EMBL/DDBJ
A/Molecule type: DNA
A/Residues: 1-68 <SEE>
A/Cross-references: EMBL:AL096844; PIDN:CAB50878.1; GSPDB:GN00070; SCORDB:SCI28.04c
A/Experimental source: strain A3(2)
C/Genetics:
A/Gene: SCORDB:SCI28.04c

Query Match 89.7%; Score 26; DB 2; Length 68;
Best Local Similarity 80.0%; Pred. No. 1.2e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|
Db 7 TRPPR 11

RESULT 34
AE1652
hypothetical protein lin1758 [imported] - Listeria innocua (strain Clp11262)
C/Species: Listeria innocua
C/Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 27-Nov-2001
C/Accession: AE1652
R;Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloesch, D.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi, I.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A/Authors: Krefit, J.; Kuhn, M.; Kunst, F.; Kurapat, G.; Madueno, E.; Maitournam, A.; Mok, C.; Schluter, T.; Simoes, N.; Tierrez, A.; Vaquez-Boland, J.A.; Voss, H.; Wehlan
A/Title: Comparative genomics of Listeria species.
A/Reference number: AB1077; MUID:21537279; PMID:11679669
A/Accession: AE1652
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-75 <GLA>
A/Cross-references: GB:AL592022; PIDN:CAC96989.1; PID:g16414245; GSPDB:GN00178
A/Experimental source: strain Clp11262
C/Genetics:
A/Gene: lin1758

Query Match 89.7%; Score 26; DB 2; Length 75;
Best Local Similarity 80.0%; Pred. No. 1.4e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|
Db 38 TKPPK 42

RESULT 35
AD0288
conserved hypothetical protein YPO2363 [imported] - Yersinia pestis (strain CO92)
C/Species: Yersinia pestis
C/Date: 02-Nov-2001 #sequence_revision 02-Nov-2001 #text_change 02-Nov-2001
C/Accession: AD0288
R;Parkhill, J.; Wren, B.W.; Thomson, N.R.; Titball, R.W.; Holden, M.T.G.; Prentice, M.; deno-Farraga, A.M.; Chillingworth, T.; Cronin, A.; Davies, R.M.; Davis, P.; Dougan, G.; il, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.; Whitehead, S.; Barrell
Nature 413, 523-527, 2001
A/Title: Genome sequence of Yersinia pestis, the causative agent of plague.
A/Reference number: AB0001; MUID:21470413; PMID:11586360
A/Accession: AD0288
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-87 <KUR>
A/Cross-references: GB:AL590842; PIDN:CAC91168.1; PID:g15980360; GSPDB:GN00175
C/Genetics:
A/Gene: YPO2363

Biochem. Biophys. Res. Commun. 200, 1066-1071, 1994

A;Title: cDNA sequence and genomic organization of mouse secretin.

A;Reference number: JC2202; MUID:94234995; PMID:8179583

A;Accession: JC2202

A;Molecule type: mRNA

A;Residues: 1-133 <LAN>

A;Cross-references: EMBL:X73580; NID:g313710; PIDN:CAA51982.1; PID:g313711

C;Comment: This protein regulates the secretion of pancreatic juices and stimulates insulin secretion.

C;Superfamily: glucagon

C;Keywords: amidated carboxyl end; duplication; hormone; secretagogue

F;1-21/Domain: signal sequence #status predicted <SIG>

F;28-133/Product: secretin #status predicted <PRO>

F;22-58/Product: secretin #status predicted <MAT>

F;36/Modified site: amidated carboxyl end (Val) (amide in mature form from following g

Query Match 89.7%; Score 26; DB 2; Length 133;
Best Local Similarity 80.0%; Pred. No. 2.4e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|

DB 129 TRPPR 133

RESULT 39
B49530
vascular endothelial growth factor homolog A2R, 14.7K - Orf virus
C;Species: Orf virus
C;Date: 07-Apr-1994 #sequence_revision 18-Nov-1994 #text_change 08-Oct-1999
C;Accession: B49530
R;Lytle, D.J.; Fraser, K.M.; Fleming, S.B.; Mercer, A.A.; Robinson, A.J.
J. Virol. 68, 84-92, 1994
A;Title: Homologs of vascular endothelial growth factor are encoded by the poxvirus or
A;Reference number: A49530; MUID:94076465; PMID:8254780
A;Contents: NZ2
A;Accession: B49530
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-133 <LYT>
A;Cross-references: GB:S67520; NID:g456897; PIDN:AAB29220.1; PID:g456899
A;Note: sequence inconsistent with nucleotide translation
A;Note: sequence extracted from NCBI backbone (NCBIN:141420, NCBI:P:141425)

Query Match 89.7%; Score 26; DB 2; Length 133;
Best Local Similarity 80.0%; Pred. No. 2.4e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|:|:|

DB 126 TRPPR 130

RESULT 40
A29622
ribosomal protein S12, mitochondrial - fruit fly (Drosophila melanogaster)
N;Alternate names: technical knockout protein
C;Species: Drosophila melanogaster
C;Date: 31-Mar-1989 #sequence_revision 31-Mar-1989 #text_change 12-Jun-2003
C;Accession: A29622
R;Roeyden, C.S.; Piroetta, V.; Jan, L.Y.
Cell 51, 165-173, 1987
A;Title: The tko locus, site of a behavioral mutation in D. melanogaster, codes for a
A;Reference number: A29622; MUID:89027001; PMID:3117373
A;Accession: A29622
A;Molecule type: mRNA
A;Residues: 1-140 <ROY>
A;Cross-references: GB:M19494; NID:g158601; PIDN:AAA28935.1; PID:g158602
C;Genetics: lko
A;Gene: lko
A;Cross-references: FlyBase:FBgn0003714
C;Superfamily: ribosomal protein S12
C;Keywords: mitochondrial; protein biosynthesis; ribosome
F;117/Modified site: beta-methylthioaspartic acid (Asp) #status predicted

Query Match 89.7%; Score 26; DB 2; Length 140;
 Best Local Similarity 80.0%; Pred. No. 2.5e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |:|
 45 TRPPR 49

Db

RESULT 41
 T27059
 hypothetical protein Y51A2A.6 - Caenorhabditis elegans
 C;Species: Caenorhabditis elegans
 C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
 C;Accession: T27059
 R;McMurray, A.
 submitted to the EMBL Data Library, October 1998
 A;Reference number: Z20304
 A;Accession: T27059
 A;Status: preliminary; translated from GB/EMBL/DBJ
 A;Molecule type: DNA
 A;Residues: 1-140 <WIL>
 A;Cross-references: EMBL:AL032635; PIDN:CRA21601.1; GSPDB:GN00023; CESP:Y51A2A.6
 A;Experimental source: clone Y51A2A
 C;Genetics:
 A;Gene: CESP:Y51A2A.6
 A;Map position: 5
 A;Introns: 93/3; 129/1

Query Match 89.7%; Score 26; DB 2; Length 140;
 Best Local Similarity 80.0%; Pred. No. 2.5e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |:|
 68 TRPPR 72

Db

RESULT 42
 T49706
 hypothetical protein B23L21.190 [imported] - Neurospora crassa
 C;Species: Neurospora crassa
 C;Date: 02-Jun-2000 #sequence_revision 02-Jun-2000 #text_change 02-Jun-2000
 C;Accession: T49706
 R;Schulte, U.; Aign, V.; Hohnel, J.; Brandt, P.; Fartmann, B.; Holland, R.; Nyakatura,
 submitted to the Protein Sequence Database, May 2000
 A;Reference number: Z25022
 A;Accession: T49706
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-144 <SCH>
 A;Cross-references: EMBL:AL356172; GSPDB:GN00116; NCSP:B23L21.190
 A;Experimental source: BAC clone B23L21; strain OR74A
 C;Genetics:
 A;Gene: NCSP:B23L21.190
 A;Map position: 6

Query Match 89.7%; Score 26; DB 2; Length 144;
 Best Local Similarity 80.0%; Pred. No. 2.6e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |:|
 130 TRPPR 134

Db

RESULT 43
 A87466
 hypothetical protein C01749 [imported] - Caulobacter crescentus
 C;Species: Caulobacter crescentus
 C;Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 20-Apr-2001
 C;Accession: A87466

R;Nierman, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, R.J.; Gwinn, M.L.; Haft, D.H.; Koln, J.; Ermolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter, J.C.; Fraser, C.I. Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001
 A;Title: Complete Genome Sequence of Caulobacter crescentus.
 A;Reference number: A87249; MUID:21173698; PMID:11259647
 A;Accession: A87466
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-148 <STO>
 A;Cross-references: GB:AE005673; NID:gl3423171; PIDN:AAK21725.1; GSPDB:GN00148
 C;Genetics:
 A;Gene: C01749

Query Match 89.7%; Score 26; DB 2; Length 148;
 Best Local Similarity 80.0%; Pred. No. 2.7e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |:|
 82 TRPPR 86

Db

RESULT 44
 S75561
 hypothetical protein sll0812 - Synecocystis sp. (strain PCC 6803)
 C;Species: Synecocystis sp.
 A;Variety: PCC 6803
 C;Date: 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change 20-Jun-2000
 C;Accession: S75561
 R;Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.; Nakamura, Y.; Miyajima, N. K.; Okumura, S.; Shimpo, S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.; Yasu DNA Res. 3, 109-136, 1996
 A;Title: Sequence analysis of the genome of the unicellular cyanobacterium Synecocyst: s.
 A;Reference number: S74322; MUID:97061201; PMID:8905231
 A;Accession: S75561
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-153 <KAN>
 A;Cross-references: EMBL:D90911; GB:AB001339; NID:gl653083; PIDN:BAA18122.1; PID:gl653;
 A;Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996
 C;Superfamily: Synecocystis hypothetical protein sll0812

Query Match 89.7%; Score 26; DB 2; Length 153;
 Best Local Similarity 80.0%; Pred. No. 2.8e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |:|
 15 TRPPK 19

Db

RESULT 45
 A69267
 transposase homolog - Archaeoglobus fulgidus
 C;Species: Archaeoglobus fulgidus
 C;Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 21-Jul-2000
 C;Accession: A69267
 R;Klenk, H.P.; Clayton, R.A.; Tomb, J.F.; White, O.; Nelson, K.E.; Ketchum, K.A.; Dodson, F.; Fleischmann, R.D.; Quackenbush, J.; Lee, N.H.; Sutton, G.G.; Gill, S.; Kirkness, E.I. Glodek, A.; Zhou, L.; Overbeek, R.; Gocayne, J.D.; Weidman, J.F.; McDonald, L. Nature 390, 364-370, 1997
 A;Authors: Uterback, T.; Cotton, M.D.; Spriggs, T.; Artiaich, P.; Kaine, B.P.; Sykes, S. Smith, H.O.; Woese, C.R.; Venter, J.C.
 A;Title: The complete genome sequence of the hyperthermophilic, sulfate-reducing archae
 A;Reference number: A69250; MUID:98049343; PMID:9389475
 A;Accession: A69267
 A;Status: preliminary; nucleic acid sequence not shown; translation not shown
 A;Molecule type: DNA
 A;Residues: 1-154 <KLB>
 A;Cross-references: GB:AE001097; GB:AE000782; NID:g2689420; PIDN:AA91094.1; PID:g26505

Query Match 89.7%; Score 26; DB 2; Length 154;
 Best Local Similarity 80.0%; Pred. No. 2.8e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 24 TRPPK 28

Db

RESULT 46
 T49798
 conserved hypothetical protein [imported] - Neurospora crassa (fragment)
 N;Alternate names: protein B9J10.350
 C;Species: Neurospora crassa
 C;Date: 02-Jun-2000 #sequence_revision 02-Jun-2000 #text_change 02-Jun-2000
 C;Accession: T49798
 R;Schulte, U.; Aign, V.; Hohisel, J.; Brandt, P.; Fartmann, B.; Holland, R.; Nyakatura,
 submitted to the Protein Sequence Database, May 2000
 A;Reference number: 225022
 A;Accession: T49798
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-163 <SCH>
 A;Cross-references: EMBL:AL356324; GSPDB:GN00116; NCSP:B9J10.350
 A;Experimental source: SAC clone B9J10; strain OR74A
 C;Genetics:
 A;Gene: NCSP:B9J10.350
 A;Map position: 6

Query Match 89.7%; Score 26; DB 2; Length 163;
 Best Local Similarity 80.0%; Pred. No. 3e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 45 TRPPR 49

Db

RESULT 47
 AB1974
 hypothetical protein alr1341 [imported] - Nostoc sp. (strain PCC 7120)
 C;Species: Nostoc sp. PCC 7120
 A;Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120
 C;Date: 14-Dec-2001 #sequence_revision 14-Dec-2001 #text_change 09-Dec-2002
 C;Accession: AB1974
 R;Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasamoto, S.; Watanabe, A.; Itriguchi,
 Nakazaki, N.; Shimpo, S.; Sugimoto, M.; Takazawa, M.; Yamada, M.; Tabata, S.
 DNA Res. 8, 205-213, 2001
 A;Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium Ana
 A;Reference number: AB1807; MUID:21595285; PMID:11759940
 A;Accession: AB1974
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-179 <KUR>
 A;Cross-references: GB:BA000019; PIDN:BA073298.1; PID:g17130688; GSPDB:GN00179
 A;Experimental source: strain PCC 7120
 C;Genetics:
 A;Gene: alr1341

Query Match 89.7%; Score 26; DB 2; Length 179;
 Best Local Similarity 80.0%; Pred. No. 3.3e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 72 TRPPK 76

Db

RESULT 48
 H83561
 hypothetical protein PA0671 [imported] - Pseudomonas aeruginosa (strain PAO1)
 C;Species: Pseudomonas aeruginosa
 C;Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000

C;Accession: H83561
 R;Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.; Hickey, M.J.; E
 adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; Lin
 ; Lory, S.; Olson, M.V.
 Nature 406, 959-964, 2000
 A;Title: Complete genome sequence of Pseudomonas aeruginosa PAO1, an opportunistic pat
 A;Reference number: A82950; MUID:20437337; PMID:10984043
 A;Accession: H83561
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-183 <STO>
 A;Cross-references: GB:AE004502; GB:AE004091; NID:G9946547; PIDN:AA04060.1; GSPDB:GN0
 A;Experimental source: strain PAO1
 C;Genetics:
 A;Gene: PA0671

Query Match 89.7%; Score 26; DB 2; Length 183;
 Best Local Similarity 80.0%; Pred. No. 3.3e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 157 TRPPR 161

Db

RESULT 49
 S45627
 acidic endoprotease precursor - Myxococcus xanthus (strain DK101)
 C;Species: Myxococcus xanthus
 A;Variety: strain DK101
 C;Date: 10-Dec-1994 #sequence_revision 26-Apr-1996 #text_change 08-Oct-1999
 C;Accession: S45627; S62857
 R;Lucas, N.; Mazaud-Aujard, C.; Brenaud, L.; Cenatiempo, Y.; Julien, R.
 Eur. J. Biochem. 222, 247-254, 1994
 A;Title: Protein purification, gene cloning and sequencing of an acidic endoprotease f
 A;Reference number: S45627; MUID:94291618; PMID:8020464
 A;Accession: S45627
 A;Molecule type: DNA
 A;Residues: 1-195 <LUC>
 A;Cross-references: EMBL:X75892; NID:G516391; PIDN:CAA53499.1; PID:G516392
 A;Experimental source: strain DK101
 A;Accession: S62857
 A;Molecule type: protein
 A;Residues: 65-101 <LUW>
 C;Genetics:
 A;Gene: Maep
 E;1-29/Domain: signal sequence #status predicted <SIG>
 E;30-64/Domain: propeptide #status predicted <PRO>
 F;65-195/Product: acidic endoprotease #status experimental <MAT>

Query Match 89.7%; Score 26; DB 2; Length 195;
 Best Local Similarity 80.0%; Pred. No. 3.5e+02;
 Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 148 TRPPR 152

Db

RESULT 50
 FOAD72
 major core protein VII precursor - human adenovirus 2
 C;Species: Mastadenovirus h2 (human adenovirus 2)
 A;Note: host Homo sapiens (man)
 C;Date: 18-Apr-1984 #sequence_revision 31-Dec-1993 #text_change 16-Jul-1999
 C;Accession: C03837; A03836
 R;Alestrom, P.; Akusjaervi, G.; Lager, M.; Yeh-kai, L.; Pettersson, U.
 J. Biol. Chem. 259, 13980-13985, 1984
 A;Title: Genes encoding the core proteins of adenovirus type 2.
 A;Reference number: A03837; MUID:85054835; PMID:6094534
 A;Accession: C03837
 A;Molecule type: DNA
 A;Residues: 1-198 <ALE>

A;Cross-references: GB:J01917; NID:G209811; PIDN:AAA92212.1; PID:G209828
R;Sung, M.T.; Cao, T.M.; Coleman, R.T.; Budelier, K.A.
Proc. Natl. Acad. Sci. U.S.A. 80, 2902-2906, 1983
A;Title: Gene and protein sequences of adenovirus protein VII, a hybrid basic chromosome
A;Reference number: A03836; MUID:83221511; PMID:6574459
A;Accession: A03836
A;Molecule type: DNA
A;Residues: 1-111/113-198 <SUN>
A;Cross-references: GB:J01917
C;Genetics:
A;Map position: 43-45
C;Superfamily: adenovirus major core protein VII
C;Keywords: core protein; late protein
F;1-24/Domain: signal sequence #status predicted <SIG>
F;25-198/Product: major core protein VII #status predicted <MCP>

Query Match 89.7%; Score 26; DB 1; Length 198;
Best Local Similarity 80.0%; Pred. No. 3.6e+02;
Matches 4; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TKPR 5
Db 193 TRPR 197

Search completed: March 3, 2004, 12:19:05
Job time : 23 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: March 3, 2004, 12:16:13 ; Search time 23 Seconds
(without alignments)
11.223 Million cell updates/sec

Title: US-09-871-974-2

Perfect score: 29

Sequence: 1 TKPPR 5

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database : Issued Patents AA:*
1: /cgn2_6/ptodata/2/iaa/5A COMB pep.*
2: /cgn2_6/ptodata/2/iaa/5B COMB pep.*
3: /cgn2_6/ptodata/2/iaa/6A COMB pep.*
4: /cgn2_6/ptodata/2/iaa/6B COMB pep.*
5: /cgn2_6/ptodata/2/iaa/PCITUS COMB pep.*
6: /cgn2_6/ptodata/2/iaa/backfiles1 pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	29	100.0	5	1	US-08-171-737-2
2	29	100.0	5	1	US-08-202-178-2
3	29	100.0	5	1	US-08-299-636-37
4	29	100.0	5	1	US-08-279-155-36
5	29	100.0	5	1	US-08-713-484-2
6	29	100.0	5	1	US-08-703-988A-36
7	29	100.0	5	1	US-08-532-294-1
8	29	100.0	5	1	US-08-454-859-1
9	29	100.0	5	2	US-08-955-263-2
10	29	100.0	5	2	US-08-612-842-36
11	29	100.0	6	1	US-08-171-737-18
12	29	100.0	6	1	US-08-299-636-38
13	29	100.0	6	1	US-08-279-155-37
14	29	100.0	6	1	US-08-703-988A-37
15	29	100.0	6	2	US-08-612-842-37
16	29	100.0	8	1	US-08-202-178-4
17	29	100.0	8	1	US-08-713-484-4
18	29	100.0	8	1	US-08-532-294-2
19	29	100.0	8	2	US-08-955-263-4
20	29	100.0	8	2	US-08-612-842-39
21	29	100.0	9	1	US-08-171-737-9
22	29	100.0	9	1	US-08-279-155-1
23	29	100.0	9	1	US-08-279-155-38
24	29	100.0	9	1	US-08-703-988A-1
25	29	100.0	9	1	US-08-703-988A-38
26	29	100.0	9	1	US-08-532-294-3
27	29	100.0	9	1	US-08-454-859-2

28	29	100.0	9	2	US-08-613-842-1	Sequence 1, Appl
29	29	100.0	9	2	US-08-613-842-38	Sequence 38, Appl
30	29	100.0	9	4	US-08-997-802-10	Sequence 10, Appl
31	29	100.0	9	4	US-08-997-802-11	Sequence 11, Appl
32	29	100.0	10	1	US-08-202-178-5	Sequence 5, Appl
33	29	100.0	10	1	US-08-713-484-5	Sequence 5, Appl
34	29	100.0	10	2	US-08-955-263-5	Sequence 5, Appl
35	29	100.0	11	1	US-08-202-178-7	Sequence 7, Appl
36	29	100.0	11	1	US-08-713-484-7	Sequence 7, Appl
37	29	100.0	11	2	US-08-955-263-7	Sequence 7, Appl
38	29	100.0	11	4	US-09-387-715-51	Sequence 51, Appl
39	29	100.0	11	4	US-09-387-715-52	Sequence 52, Appl
40	29	100.0	12	1	US-08-299-636-39	Sequence 39, Appl
41	29	100.0	60	4	US-09-381-546-13	Sequence 13, Appl
42	29	100.0	98	4	US-09-252-991A-25421	Sequence 25421, A
43	29	100.0	136	4	US-09-252-991A-21156	Sequence 21156, A
44	29	100.0	420	4	US-09-489-039A-8820	Sequence 8820, Ap
45	29	100.0	576	4	US-09-367-206-1	Sequence 1, Appl
46	29	100.0	576	4	US-09-367-206-21	Sequence 21, Appl
47	29	100.0	576	4	US-09-367-206-22	Sequence 22, Appl
48	29	100.0	576	4	US-09-367-206-23	Sequence 23, Appl
49	29	100.0	892	3	US-08-857-076-42	Sequence 42, Appl
50	29	100.0	1235	1	US-08-118-101A-2	Sequence 2, Appl
51	26	89.7	14	4	US-09-608-285A-44	Sequence 44, Appl
52	26	89.7	14	4	US-09-557-800C-44	Sequence 44, Appl
53	26	89.7	29	4	US-09-287-849-30	Sequence 30, Appl
54	26	89.7	44	4	US-09-645-470-14	Sequence 14, Appl
55	26	89.7	67	4	US-09-107-532A-3781	Sequence 3781, Ap
56	26	89.7	128	4	US-09-252-991A-28839	Sequence 28839, A
57	26	89.7	132	4	US-09-125-642C-15	Sequence 15, Appl
58	26	89.7	132	4	US-09-431-888-11	Sequence 11, Appl
59	26	89.7	133	4	US-09-431-888-2	Sequence 2, Appl
60	26	89.7	140	4	US-09-732-210-1700	Sequence 1700, Ap
61	26	89.7	147	4	US-09-252-991A-18713	Sequence 18713, A
62	26	89.7	178	4	US-09-252-991A-18672	Sequence 18672, A
63	26	89.7	211	4	US-09-252-991A-36873	Sequence 36873, A
64	26	89.7	211	4	US-09-252-991A-30029	Sequence 30029, A
65	26	89.7	220	4	US-09-252-991A-16739	Sequence 16739, A
66	26	89.7	234	4	US-09-252-991A-28621	Sequence 28621, A
67	26	89.7	245	4	US-09-252-991A-32366	Sequence 32366, A
68	26	89.7	253	4	US-09-252-991A-20329	Sequence 20329, A
69	26	89.7	263	1	US-08-565-386-9	Sequence 9, Appl
70	26	89.7	264	4	US-09-252-991A-22289	Sequence 22289, A
71	26	89.7	282	4	US-09-252-991A-19978	Sequence 19978, A
72	26	89.7	290	4	US-09-564-951B-4	Sequence 4, Appl
73	26	89.7	304	4	US-09-355-040-13	Sequence 13, Appl
74	26	89.7	307	4	US-09-355-040-14	Sequence 14, Appl
75	26	89.7	308	4	US-09-050-739-94	Sequence 94, Appl
76	26	89.7	309	4	US-09-252-991A-17744	Sequence 17744, A
77	26	89.7	313	4	US-09-540-236-2724	Sequence 2724, Ap
78	26	89.7	314	3	US-08-612-973-42	Sequence 42, Appl
79	26	89.7	314	3	US-08-927-597-42	Sequence 42, Appl
80	26	89.7	316	4	US-09-252-991A-30881	Sequence 30881, A
81	26	89.7	319	3	US-08-612-973-44	Sequence 44, Appl
82	26	89.7	319	3	US-08-927-597-44	Sequence 44, Appl
83	26	89.7	321	4	US-09-252-991A-24925	Sequence 24925, A
84	26	89.7	332	4	US-09-543-681A-5632	Sequence 5632, Ap
85	26	89.7	338	3	US-08-612-973-38	Sequence 38, Appl
86	26	89.7	338	3	US-08-927-597-38	Sequence 38, Appl
87	26	89.7	343	3	US-08-612-973-40	Sequence 40, Appl
88	26	89.7	343	3	US-08-927-597-40	Sequence 40, Appl
89	26	89.7	349	4	US-09-252-991A-30578	Sequence 30578, A
90	26	89.7	350	4	US-09-818-780-95	Sequence 95, Appl
91	26	89.7	359	4	US-09-269-137-8	Sequence 8, Appl
92	26	89.7	359	4	US-09-355-040-30	Sequence 30, Appl
93	26	89.7	375	4	US-09-540-236-2085	Sequence 2085, Ap
94	26	89.7	384	4	US-09-252-991A-18786	Sequence 18786, A
95	26	89.7	393	4	US-09-107-532A-3712	Sequence 3712, Ap
96	26	89.7	395	3	US-08-928-442-1	Sequence 1, Appl
97	26	89.7	428	3	US-08-570-157-5	Sequence 5, Appl
98	26	89.7	428	3	US-08-029-170-31	Sequence 31, Appl
99	26	89.7	428	4	US-09-076-510-5	Sequence 5, Appl
100	26	89.7	428	4	US-09-004-349-5	Sequence 5, Appl

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
Db 1 TKPR 5

RESULT 3
US-08-299-636-37
Sequence 37, Application US/08299636
Patent No. 5659041
GENERAL INFORMATION:
APPLICANT: POLLAK, Alfred
APPLICANT: KIRBY, Robert A.
APPLICANT: DUNN-DUFAULT, Robert
TITLE OF INVENTION: HYDRAZINO-TYPE RADIONUCLIDE CHELATORS
NUMBER OF SEQUENCES: 39
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/299,636
FILING DATE: 02-SEP-1994
CLASSIFICATION: 534
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/092,911
FILING DATE: 18-JUL-1993
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/262/ALLE
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 37:
SEQUENCE CHARACTERISTICS:
LENGTH: 5 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-299-636-37

Query Match 100.0%; Score 29; DB 1; Length 5;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
Db 1 TKPR 5

RESULT 4
US-08-279-155-36
Sequence 36, Application US/08279155
Patent No. 5662885
GENERAL INFORMATION:
APPLICANT: POLLAK, Alfred
APPLICANT: GOODEBY, Anne
TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
NUMBER OF SEQUENCES: 38
CORRESPONDENCE ADDRESS:
ADDRESSEE: NIKAIKO, MARCELSTEIN, MURRAY & ORAM LLP

STREET: 655 Fifteenth Street, N. W., Suite 330 - G
STREET: Street Lobby
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-5701
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/279,155
FILING DATE: 22-JUL-1994
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: MURRAY, Robert B.
REGISTRATION NUMBER: 22,980
REFERENCE/DOCKET NUMBER: P8074-4005
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202/638-5000
TELEFAX: 202/638-4810
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 5 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-279-155-36

Query Match 100.0%; Score 29; DB 1; Length 5;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
Db 1 TKPR 5

RESULT 5
US-08-713-484-2
Sequence 2, Application US/08713484
Patent No. 5679642
GENERAL INFORMATION:
APPLICANT: Goodbody, Anne
APPLICANT: POLLAK, Alfred
TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/713,484
FILING DATE: 13-SEP-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/202,178
FILING DATE: 25-FEB-1994
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/290/ALLE
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399

Qy 1 TKPPR 5

STATE: DC
COUNTRY: USA
ZIP: 20005-5701
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: IBM PC compatible
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/454,859
FILING DATE: 31-MAY-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Berman, Richard J.
REGISTRATION NUMBER: 39,107
REFERENCE/DOCKET NUMBER: 8074-5007
TELEPHONE: (202) 638-5000
TELEFAX: (202) 638-4810
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 5 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-454-859-1

Query Match 100.0%; Score 29; DB 1; Length 5;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 1 TKPPR 5

RESULT 9
US-08-955-263-2
Sequence 2, Application US/08955263
Patent No. 5866544
GENERAL INFORMATION:
APPLICANT: Goodbody, Anne
APPLICANT: Pollak, Alfred
TITLE OF INVENTION: PEPTIDE-CHLATOR CONJUGATES
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/955,263
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/713,484
FILING DATE: 13-SEP-1996
APPLICATION NUMBER: US 08/202,178
FILING DATE: 25-FEB-1994
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/290/ALLE
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399
TELEX: 904136

INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 5 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-955-263-2

Query Match 100.0%; Score 29; DB 2; Length 5;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 1 TKPPR 5

RESULT 10
US-08-612-842-36
Sequence 36, Application US/08612842
Patent No. 5976495
GENERAL INFORMATION:
APPLICANT: POLLAK, ALFRED
APPLICANT: GOODBODY, ANNE
TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
NUMBER OF SEQUENCES: 39
CORRESPONDENCE ADDRESS:
ADDRESSEE: NIKAIKO, MARCELSTEIN, MURRAY & ORAM
STREET: 655 15TH STREET, NW, G STREET LOBBY, SUITE
STREET: 330
CITY: WASHINGTON
STATE: DC
COUNTRY: USA
ZIP: 20005-5701
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/612,842
FILING DATE: 20-MAR-1996
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: BERMAN, RICHARD J.
REGISTRATION NUMBER: 39107
REFERENCE/DOCKET NUMBER: 8012-6002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202 638-5000
TELEFAX: 202 638-4810
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 5 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-612-842-36

Query Match 100.0%; Score 29; DB 2; Length 5;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 1 TKPPR 5

RESULT 11
US-08-171-737-18
Sequence 18, Application US/08171737
Patent No. 5480970
GENERAL INFORMATION:

APPLICANT: Pollak, Alfred
APPLICANT: Goodbody, Anne
TITLE OF INVENTION: METAL CHELATORS
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/171,737
FILING DATE: 22-DEC-1993
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/253/ALLE
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 6 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-171-737-18

Query Match 100.0%; Score 29; DB 1; Length 6;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 2 TKPPR 6

RESULT 12
US-08-299-636-38
; Sequence 38, Application US/08299636
; Patent No. 5659041
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: KIRBY, Robert A.
; APPLICANT: DUNN-DUFFAULT, Robert
; TITLE OF INVENTION: HYDRAZINO-TYPE RADIONUCLIDE CHELATORS
; TITLE OF INVENTION: HAVING AN N3S CONFIGURATION
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: 3000 K Street, N.W., Suite 500
; CITY: Washington
; STATE: D.C.
; COUNTRY: USA
; ZIP: 20007-5109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/299,636
; FILING DATE: 02-SEP-1994
; CLASSIFICATION: 534
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/092,911

FILING DATE: 18-JUL-1993
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/262/ALLE
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 38:
SEQUENCE CHARACTERISTICS:
LENGTH: 6 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
US-08-299-636-38

Query Match 100.0%; Score 29; DB 1; Length 6;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 2 TKPPR 6

RESULT 13
US-08-279-155-37
; Sequence 37, Application US/08279155
; Patent No. 5662885
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: GOODBODY, Anne
; TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NAKAIDO, MARCELSTEIN, MURRAY & ORAM LLP
; STREET: 655 Fifteenth Street, N. W., Suite 330 - G
; STREET: Street Lobby
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/279,155
; FILING DATE: 22-JUL-1994
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: MURRAY, Robert B.
; REGISTRATION NUMBER: 22,980
; REFERENCE/DOCKET NUMBER: P8074-4005
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202/638-5000
; TELEFAX: 202/638-4810
; INFORMATION FOR SEQ ID NO: 37:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 6
; OTHER INFORMATION: /note= "Arg at position 6 has an OH
; OTHER INFORMATION: group."
; US-08-279-155-37

Query Match 100.0%; Score 29; DB 1; Length 6;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
Db 2 TKPR 6

RESULT 14

US-08-703-988A-37
; Sequence 37, Application US/08703988A
; Patent No. 5780006
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: GOODBODY, Anne
; TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKALDO, MARCELSTEIN, MURRAY & ORAM
; STREET: 655 Fifteenth Street, N. W., Suite 330
; STREET: -
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/703,988A
; FILING DATE: 28-AUG-1996
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/279,155
; FILING DATE: 22-JUL-1994
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: MURRAY, Robert B.
; REGISTRATION NUMBER: 22,980
; REFERENCE/DOCKET NUMBER: P8074-6011
; TELEPHONE: 202/638-5000
; TELEFAX: 202/638-4810
; INFORMATION FOR SEQ ID NO: 37:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 6
; OTHER INFORMATION: /note= "Arg at position 6 has
; OTHER INFORMATION: an OH group."
US-08-703-988A-37

Query Match 100.0%; Score 29; DB 1; Length 6;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
Db 2 TKPR 6

RESULT 15

US-08-612-842-37
; Sequence 37, Application US/08612842
; Patent No. 5976455
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: GOODBODY, Anne
; TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKALDO, MARCELSTEIN, MURRAY & ORAM
; STREET: 655 15TH STREET, NW, G STREET LOBBY, SUITE
; STREET: 330
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/612,842
; FILING DATE: 20-MAR-1996
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: BERMAN, RICHARD J.
; REGISTRATION NUMBER: 39107
; REFERENCE/DOCKET NUMBER: 8012-6002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202 638-5000
; TELEFAX: 202 638-4810
; INFORMATION FOR SEQ ID NO: 37:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 6
; OTHER INFORMATION: /note= "Position 6 has an -OH
; OTHER INFORMATION: substituent."
US-08-612-842-37

Query Match 100.0%; Score 29; DB 2; Length 6;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
Db 2 TKPR 6

RESULT 16

US-08-202-178-4
; Sequence 4, Application US/08202178
; Patent No. 5569745
; GENERAL INFORMATION:
; APPLICANT: Goodbody, Anne
; APPLICANT: Pollak, Alfred
; TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Hardner
; STREET: Suite 500, 3000 K Street, N.W.
; CITY: Washington, D.C.
; COUNTRY: USA
; ZIP: 20007-5109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/202,178
FILING DATE: 25-FEB-1994
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/258/ALLE
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 8 amino acids
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Ser substituted with
OTHER INFORMATION: picolinic acid (Pic)."
FEATURE:
NAME/KEY: Modified-site
LOCATION: 2
OTHER INFORMATION: /note= "Cys substituted with
OTHER INFORMATION: acetamidomethyl (AcM)."
US-08-202-178-4
Query Match 100.0%; Score 29; DB 1; Length 8;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
DB 4 TKPPR 8
RESULT 17
US-08-713-484-4
Sequence 4, Application US/08713484
Patent No. 5679642
GENERAL INFORMATION:
APPLICANT: Goodbody, Anne
TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/713,484
FILING DATE: 13-SEP-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/202,178
FILING DATE: 25-FEB-1994
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/250/ALLE
TELEPHONE: (202) 672-5300

TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 8 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Ser substituted with
OTHER INFORMATION: picolinic acid (Pic)."
FEATURE:
NAME/KEY: Modified-site
LOCATION: 2
OTHER INFORMATION: /note= "Cys substituted with
OTHER INFORMATION: acetamidomethyl (AcM)."
US-08-713-484-4
Query Match 100.0%; Score 29; DB 1; Length 8;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
DB 4 TKPPR 8
RESULT 18
US-08-592-294-2
Sequence 2, Application US/08592294
Patent No. 5789555
GENERAL INFORMATION:
APPLICANT: POLLAK, ALFRED
TITLE OF INVENTION: IMMOBILIZED LABELLING METHOD
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: NIKALDO, MARCELSTEIN, MURRAY & ORAM LLP
STREET: 655 FIFTEENTH ST., N.W., SUITE 330-G STREET
CITY: WASHINGTON
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20005-5701
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/592,294
FILING DATE:
CLASSIFICATION: 534
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/CA94/00637
FILING DATE: 16-NOV-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/152,680
FILING DATE: 16-NOV-1993
ATTORNEY/AGENT INFORMATION:
NAME: MURRAY, ROBERT B.
REGISTRATION NUMBER: 22,980
REFERENCE/DOCKET NUMBER: P8012-5002
TELEPHONE: 202-638-5000
TELEFAX: 202-638-4810
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 8 amino acids
TYPE: amino acid
STRANDEDNESS: single

TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1
OTHER INFORMATION: /note= "Picolinic acid is attached to Ser of position 1."
US-08-592-294-2

Query Match 100.0%; Score 29; DB 1; Length 8;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 4 TKPPR 8

RESULT 19
US-08-955-263-4
; Sequence 4, Application US/08955263
; Patent No. 5866544
; GENERAL INFORMATION:
; APPLICANT: Goodbody, Anne
; APPLICANT: Pollak, Alfred
; TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: Suite 500, 3000 K Street, N.W.
; CITY: Washington, D.C.
; COUNTRY: USA
; ZIP: 20007-5109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/955,263
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/713,484
; FILING DATE: 13-SEP-1996
; APPLICATION NUMBER: US 08/202,178
; FILING DATE: 25-FEB-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Bent, Stephen A.
; REGISTRATION NUMBER: 29,768
; REFERENCE/DOCKET NUMBER: 16777/290/ALLE
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 672-5300
; TELEFAX: (202) 672-5399
; TELEX: 904136
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 8 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "Ser substituted with picolinic acid (Pic)."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 2
; OTHER INFORMATION: /note= "Cys substituted with acetamidomethyl (Acm)."
US-08-955-263-4

Query Match 100.0%; Score 29; DB 2; Length 8;

Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 4 TKPPR 8

RESULT 20
US-08-612-842-39
; Sequence 39, Application US/08612842
; Patent No. 5976495
; GENERAL INFORMATION:
; APPLICANT: POLLAK, ALFRED
; APPLICANT: GOODBODY, ANNE
; TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKAI, DO, MARCELSTEIN, MURRAY & ORAM
; STREET: 655 15TH STREET, NW, G STREET LOBBY, SUITE
; STREET: 330
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/612,842
; FILING DATE: 20-MAR-1996
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: BERMAN, RICHARD J
; REGISTRATION NUMBER: 39107
; REFERENCE/DOCKET NUMBER: 8012-6002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202 638-5000
; TELEFAX: 202 638-4810
; INFORMATION FOR SEQ ID NO: 39:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 8 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "Position 1 has a sarcosine substituent."
US-08-612-842-39

Query Match 100.0%; Score 29; DB 2; Length 8;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 4 TKPPR 8

RESULT 21
US-08-171-737-9
; Sequence 9, Application US/08171737
; Patent No. 5480970
; GENERAL INFORMATION:
; APPLICANT: POLLAK, ALFRED
; APPLICANT: GOODBODY, ANNE
; TITLE OF INVENTION: METAL CHELATORS
; NUMBER OF SEQUENCES: 19

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: Suite 500, 3000 K Street N.W.
; CITY: Washington, D.C.
; COUNTRY: USA
; ZIP: 20007-5109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/171,737
; FILING DATE: 22-DEC-1993
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Bent, Stephen A.
; REGISTRATION NUMBER: 29,768
; REFERENCE/DOCKET NUMBER: 16777/253/ALLE
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 672-5300
; TELEFAX: (202) 672-5399
; TELEX: 804136
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "X at position 1 is
; OTHER INFORMATION: picolinic acid (Pic)."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 3
; OTHER INFORMATION: /note= "Cys at position 3 is
; OTHER INFORMATION: substituted with acetamidomethyl (AcM)."
; US-08-171-737-9

Query Match 100.0%; Score 29; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 5 TKPPR 9

RESULT 22
US-08-279-155-1
; Sequence 1, Application US/08279155
; Patent No. 5662885
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: GOODBODY, Anne
; TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKAI, DO, MARCELSTEIN, MURRAY & ORAM LLP
; STREET: 655 Fifteenth Street, N. W., Suite 330 - G
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/279,155
; FILING DATE: 22-JUL-1994
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: MURRAY, Robert B.
; REGISTRATION NUMBER: 22,980
; REFERENCE/DOCKET NUMBER: P8074-4005
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202/638-5000
; TELEFAX: 202/638-4810
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "Gly at position 1 has an
; OTHER INFORMATION: N,N-dimethyl group."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 3
; OTHER INFORMATION: /note= "Cys at position 3 has an
; OTHER INFORMATION: AcM group."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 9
; OTHER INFORMATION: /note= "Arg at position 10 is
; OTHER INFORMATION: unsubstituted or has an OH group."
; US-08-279-155-1

Query Match 100.0%; Score 29; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 5 TKPPR 9

RESULT 23
US-08-279-155-38
; Sequence 38, Application US/08279155
; Patent No. 5663885
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: GOODBODY, Anne
; TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKAI, DO, MARCELSTEIN, MURRAY & ORAM LLP
; STREET: 655 Fifteenth Street, N. W., Suite 330 - G
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/279,155
; FILING DATE: 22-JUL-1994
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: MURRAY, Robert B.
; REGISTRATION NUMBER: 22,980
; REFERENCE/DOCKET NUMBER: P8074-4005

```

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202/638-5000
; TELEFAX: 202/638-4810
; INFORMATION FOR SEQ ID NO: 38:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "Gly at position 1 has an
; OTHER INFORMATION: N,N-dimethyl group."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 3
; OTHER INFORMATION: /note= "Cys at position 3 has an
; OTHER INFORMATION: Acn group."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 9
; OTHER INFORMATION: /note= "Arg at position 9 is
; OTHER INFORMATION: unsubstituted or has an OH group."
;
US-08-279-155-38

```

```

Query Match 100.0%; Score 29; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 1 TKPPR 5
Db 5 TKPPR 9

```

```

RESULT 24
US-08-703-988A-1
; Sequence 1, Application US/08703988A
; Patent No. 5780006
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: GOODBODY, Anne
; TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE
; TITLE OF INVENTION: CHELATORS
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKAIKO, MARCELSTEIN, MURRAY & ORAM
; ADDRESSEE: LLP
; STREET: 655 Fifteenth Street, N. W., Suite 330
; STREET:
; STREET: G Street Lobby
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/703,988A
; FILING DATE: 28-AUG-1996
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/279,155
; FILING DATE: 22-JUL-1994
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: MURRAY, Robert B.
; REGISTRATION NUMBER: 22,980
; REFERENCE/DOCKET NUMBER: P8074-6011

```

```

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202/638-5000
; TELEFAX: 202/638-4810
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "Gly at position 1 has
; OTHER INFORMATION: an N,N-dimethyl group."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 3
; OTHER INFORMATION: /note= "Cys at position 3 has
; OTHER INFORMATION: an Acn group."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 9
; OTHER INFORMATION: /note= "Arg at position 9 is
; OTHER INFORMATION: unsubstituted or has an OH group."
;
US-08-703-988A-1

```

```

Query Match 100.0%; Score 29; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 1 TKPPR 5
Db 5 TKPPR 9

```

```

RESULT 25
US-08-703-988A-38
; Sequence 38, Application US/08703988A
; Patent No. 5780006
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: GOODBODY, Anne
; TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE
; TITLE OF INVENTION: CHELATORS
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKAIKO, MARCELSTEIN, MURRAY & ORAM
; ADDRESSEE: LLP
; STREET: 655 Fifteenth Street, N. W., Suite 330
; STREET:
; STREET: G Street Lobby
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/703,988A
; FILING DATE: 28-AUG-1996
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/279,155
; FILING DATE: 22-JUL-1994
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: MURRAY, Robert B.
; REGISTRATION NUMBER: 22,980
; REFERENCE/DOCKET NUMBER: P8074-6011

```

```

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202/638-5000
; TELEFAX: 202/638-4810
; INFORMATION FOR SEQ ID NO: 38:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "gly at position 1 has
; OTHER INFORMATION: an N,N-dimethyl group."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 3
; OTHER INFORMATION: /note= "Cys at position 3 has
; OTHER INFORMATION: an AcM group."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 9
; OTHER INFORMATION: /note= "Arg at position 9 is
; OTHER INFORMATION: unsubstituted or has an OH group."
US-08-703-988A-38

```

```

Query Match 100.0%; Score 29; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy 1 TKPPR 5
Db 5 TKPPR 9

```

RESULT 26

```

US-08-592-294-3
; Sequence 3, Application US/08592294
; Patent No. 5789555
; GENERAL INFORMATION:
; APPLICANT: POLLAK, ALFRED
; TITLE OF INVENTION: IMMOBILIZED LABELLING METHOD
; NUMBER OF SEQUENCES: 3
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKAIKO, MARCELSTEIN, MURRAY & ORAM LLP
; STREET: 655 FIFTEENTH ST., N.W., SUITE 330-G STREET
; CITY: WASHINGTON
; STATE: D.C.
; COUNTRY: U.S.A.
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/592,294
; FILING DATE:
; CLASSIFICATION: 534
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/CA94/00637
; FILING DATE: 16-NOV-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/152,680
; FILING DATE: 16-NOV-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: MURRAY, ROBERT B
; REGISTRATION NUMBER: 22,980
; REFERENCE/DOCKET NUMBER: P8012-5002
; TELECOMMUNICATION INFORMATION:

```

```

; TELEPHONE: 202-638-5000
; TELEFAX: 202-638-4810
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 1
; OTHER INFORMATION: /note= "N,N'-dimethyl is attached to Gly
; OTHER INFORMATION: of position 1."
US-08-592-294-3

```

```

Query Match 100.0%; Score 29; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy 1 TKPPR 5
Db 5 TKPPR 9

```

RESULT 27

```

US-08-454-859-2
; Sequence 2, Application US/08454859
; Patent No. 580458
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; TITLE OF INVENTION: SEQUESTERED IMAGING AGENTS
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIKAIKO, Marmelstein, Murray & Oram
; STREET: 655 Fifteenth Street, NW, Suite 330, G.
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-5701
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/454,859
; FILING DATE: 31-MAY-1995
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: Bertan, Richard J.
; REGISTRATION NUMBER: 39,107
; REFERENCE/DOCKET NUMBER: 8074-5007
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 638-5000
; TELEFAX: (202) 638-4810
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "Position 1 has either a
; OTHER INFORMATION: dimethyl substituent or an N,N'-dimethyl substituent."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 9
; OTHER INFORMATION: /note= "Position 9 has an -OH

```

OTHER INFORMATION: substituent."

-454-859-2

ry Match 100.0%; Score 29; DB 1; Length 9;
t Local Similarity 100.0%; Pred. No. 3e+05;
ches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 TKPR 5
|||||
5 TKPR 9

IT 28

8-612-842-1
quence 1, Application US/08612842
tent No. 5976495
GENERAL INFORMATION:
APPLICANT: POLLAK, ALFRED
TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
NUMBER OF SEQUENCES: 39
CORRESPONDENCE ADDRESS:
ADDRESSES: NIKALDO, MARCELSTEIN, MURRAY & ORAM
STREET: 655 15TH STREET, NW, G STREET LOBBY, SUITE
CITY: WASHINGTON
STATE: DC
COUNTRY: USA
ZIP: 20005-5701
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/612.842
FILING DATE: 20-MAR-1996
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: BERMAN, RICHARD J
REGISTRATION NUMBER: 39107
REFERENCE/DOCKET NUMBER: 8012-6002
TELEPHONE: 202 638-5000
TELEFAX: 202 638-4810
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 9 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Position 1 has either an N,N-dibenzyl, an N,N-diethyl, or a sarcosine substituent."

Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 TKPR 5
|||||
5 TKPR 9

1 TKPR 5
|||||
5 TKPR 9

RESULT 29

US-08-612-842-38
Sequence 38, Application US/08612842
Patent No. 5976495
GENERAL INFORMATION:
APPLICANT: POLLAK, ALFRED
TITLE OF INVENTION: PEPTIDE DERIVED RADIONUCLIDE CHELATORS
NUMBER OF SEQUENCES: 39
CORRESPONDENCE ADDRESS:
ADDRESSES: NIKALDO, MARCELSTEIN, MURRAY & ORAM
STREET: 655 15TH STREET, NW, G STREET LOBBY, SUITE
CITY: WASHINGTON
STATE: DC
COUNTRY: USA
ZIP: 20005-5701
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/612.842
FILING DATE: 20-MAR-1996
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: BERMAN, RICHARD J
REGISTRATION NUMBER: 39107
REFERENCE/DOCKET NUMBER: 8012-6002
TELEPHONE: 202 638-5000
TELEFAX: 202 638-4810
INFORMATION FOR SEQ ID NO: 38:
SEQUENCE CHARACTERISTICS:
LENGTH: 9 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 9
OTHER INFORMATION: /note= "Position 9 is either unsubstituted or has an -OH substituent."

US-08-612-842-38

Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPR 5

DB 5 TKPR 9

RESULT 30

US-08-997-802-10
Sequence 10, Application US/08997802
Patent No. 6334996
GENERAL INFORMATION:
APPLICANT: WONG, Ernest
TITLE OF INVENTION: CHELATORS THAT PREDOMINATELY FORM A SINGLE
TITLE OF INVENTION: STEREOISOMERIC SPECIES UPON COORDINATION TO A METAL
FILE REFERENCE: 8298-7019
CURRENT APPLICATION NUMBER: US/08/997.802

US-08-997-802-10

Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPR 5

DB 5 TKPR 9

CURRENT FILING DATE: 1997-12-24
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 10

LENGTH: 9
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
NAME/KEY: MOD_RES
LOCATION: (1)_RES
OTHER INFORMATION: dimethylglycine
FEATURE:
NAME/KEY: MOD_RES
LOCATION: (2)_
OTHER INFORMATION: L-t-butylglycine
FEATURE:
NAME/KEY: MOD_RES
LOCATION: (3)_
OTHER INFORMATION: L-cysteine with an acetoamidomethyl protecting
OTHER INFORMATION: group attached via the Sulfur atom.
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Synthetic
OTHER INFORMATION: Peptide
S-08-997-802-10

Query Match 100.0%; Score 29; DB 4; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Y 1 TKPR 5
|||
Db 5 TKPR 9

RESULT 31
JS-08-997-802-11
; Sequence 11, Application US/08997802
; Patent No. 6334996
; GENERAL INFORMATION:
; APPLICANT: WONG, Ernest
; TITLE OF INVENTION: CHELATORS THAT PREDOMINATELY FORM A SINGLE
; TITLE OF INVENTION: STEREOISOMERIC SPECIES UPON COORDINATION TO A METAL
; TITLE OF INVENTION: CENTER
; FILE REFERENCE: 8298-7019
; CURRENT APPLICATION NUMBER: US/08/997,802
; CURRENT FILING DATE: 1997-12-24
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 11
; LENGTH: 9
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: MOD_RES
; LOCATION: (1)_
; OTHER INFORMATION: Dimethylglycine.
; FEATURE:
; NAME/KEY: MOD_RES
; LOCATION: (2)_
; OTHER INFORMATION: L-t-butyl glycine
; FEATURE:
; NAME/KEY: MOD_RES
; LOCATION: (3)_
; OTHER INFORMATION: L-cysteine.
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: Peptide
US-08-997-802-11

Query Match 100.0%; Score 29; DB 4; Length 9;
Best Local Similarity 100.0%; Pred. No. 3e+05;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
|||
Db 5 TKPR 9

RESULT 32
US-08-202-178-5
; Sequence 5, Application US/08202178
; Patent No. 5589745
; GENERAL INFORMATION:
; APPLICANT: Goodbody, Anne
; APPLICANT: Pollak, Alfred
; TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: Suite 500, 3000 K Street, N.W.
; CITY: Washington, D.C.
; COUNTRY: USA
; ZIP: 20007-5109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/202,178
; FILING DATE: 25-FEB-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Bent, Stephen A.
; REGISTRATION NUMBER: 29,768
; REFERENCE/DOCKET NUMBER: 16777/258/ALLE
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 672-5300
; TELEFAX: (202) 672-5399
; TELEX: 904136
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note= "Ser substituted with
; OTHER INFORMATION: picolinic acid (Pic)."
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 2
; OTHER INFORMATION: /note= "Cys substituted with
; OTHER INFORMATION: acetamidomethyl (Acm)."
US-08-202-178-5

Query Match 100.0%; Score 29; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 16;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPR 5
|||
Db 6 TKPR 10

RESULT 33
US-08-713-484-5
; Sequence 5, Application US/08713484
; Patent No. 5679642
; GENERAL INFORMATION:
; APPLICANT: Goodbody, Anne
; APPLICANT: Pollak, Alfred
; TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES

NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/713,484
FILING DATE: 13-SEP-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/202,178
FILING DATE: 25-FEB-1994
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/290/ALLE
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 10 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Ser substituted with
OTHER INFORMATION: picolinic acid (Pic)."
FEATURE:
NAME/KEY: Modified-site
LOCATION: 2
OTHER INFORMATION: /note= "Cys substituted with
OTHER INFORMATION: acetamidomethyl (Acm)."
US-08-713-484-5

Query Match 100.0%; Score 29; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 16;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TKPPR 5
Db 6 TKPPR 10

RESULT 34
US-08-955-263-5
Sequence 5, Application US/08955263
Patent No. 5866544
GENERAL INFORMATION:
APPLICANT: Goodbody, Anne
APPLICANT: Pollak, Alfred
TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/955,263
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/713,484
FILING DATE: 13-SEP-1996
APPLICATION NUMBER: US 08/202,178
FILING DATE: 25-FEB-1994
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/290/ALLE
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 10 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Ser substituted with
OTHER INFORMATION: picolinic acid (Pic)."
FEATURE:
NAME/KEY: Modified-site
LOCATION: 2
OTHER INFORMATION: /note= "Cys substituted with
OTHER INFORMATION: acetamidomethyl (Acm)."
US-08-955-263-5

Query Match 100.0%; Score 29; DB 2; Length 10;
Best Local Similarity 100.0%; Pred. No. 16;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TKPPR 5
Db 6 TKPPR 10

RESULT 35
US-08-202-178-7
Sequence 7, Application US/08202178
Patent No. 5569745
GENERAL INFORMATION:
APPLICANT: Goodbody, Anne
APPLICANT: Pollak, Alfred
TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/202,178
FILING DATE: 25-FEB-1994
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/258/ALLE
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 672-5300

TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 11 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Ser is substituted with
benzoylmercaptoacetic acid (Bz-MA)."
FEATURE:
NAME/KEY: Modified-site
LOCATION: 2
OTHER INFORMATION: /note= "Cys is substituted with
acetamidomethyl (Acm)."
US-08-202-178-7
Query Match 100.0%; Score 29; DB 1; Length 11;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPR 5
Db 7 TKPR 11
RESULT 36
US-08-713-484-7
Sequence 7, Application US/08713484
Patent No. 5679642
GENERAL INFORMATION:
APPLICANT: Goodbody, Anne
APPLICANT: Pollak, Alfred
TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/713,484
FILING DATE: 13-SEP-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/202,178
FILING DATE: 25-FEB-1994
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/290/ALLE
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 11 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Ser is substituted with
benzoylmercaptoacetic acid (Bz-MA)."
FEATURE:
NAME/KEY: Modified-site
LOCATION: 2
OTHER INFORMATION: /note= "Cys is substituted with
acetamidomethyl (Acm)."
US-08-955-263-7

TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 11 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Ser is substituted with
benzoylmercaptoacetic acid (Bz-MA)."
FEATURE:
NAME/KEY: Modified-site
LOCATION: 2
OTHER INFORMATION: /note= "Cys is substituted with
acetamidomethyl (Acm)."
US-08-713-484-7
Query Match 100.0%; Score 29; DB 1; Length 11;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPR 5
Db 7 TKPR 11
RESULT 37
US-08-955-263-7
Sequence 7, Application US/08955263
Patent No. 5866544
GENERAL INFORMATION:
APPLICANT: Goodbody, Anne
APPLICANT: Pollak, Alfred
TITLE OF INVENTION: PEPTIDE-CHELATOR CONJUGATES
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: Suite 500, 3000 K Street, N.W.
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/955,263
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/713,484
FILING DATE: 13-SEP-1996
APPLICATION NUMBER: US 08/202,178
FILING DATE: 25-FEB-1994
ATTORNEY/AGENT INFORMATION:
NAME: Bent, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 16777/290/ALLE
TELEPHONE: (202) 672-5300
TELEFAX: (202) 672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 11 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note= "Ser is substituted with
benzoylmercaptoacetic acid (Bz-MA)."
FEATURE:
NAME/KEY: Modified-site
LOCATION: 2
OTHER INFORMATION: /note= "Cys is substituted with
acetamidomethyl (Acm)."
US-08-955-263-7
Query Match 100.0%; Score 29; DB 2; Length 11;
Best Local Similarity 100.0%; Pred. No. 17;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|||||
Db 7 TKPPR 11

RESULT 38

US-09-387-715-51
; Sequence 51, Application US/09387715
; Patent No. 6551574
; GENERAL INFORMATION:
; APPLICANT: Sharma, Shubb
; TITLE OF INVENTION: Tuftsin Metallopeptides Analogs and Uses Thereof
; FILE REFERENCE: 1173/1D794US1
; CURRENT APPLICATION NUMBER: US/09/387,715
; CURRENT FILING DATE: 1999-08-30
; PRIOR APPLICATION NUMBER: PCT/US99/05693
; PRIOR FILING DATE: 1999-03-18
; NUMBER OF SEQ ID NOS: 55
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 51
; LENGTH: 11
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: synthetic
; OTHER INFORMATION: peptide
; NAME/KEY: MOD_RES
; LOCATION: 6
; OTHER INFORMATION: Ahe
; NAME/KEY: modified residue
; LOCATION: 8,10
; OTHER INFORMATION: D-amino acid
US-09-387-715-51

Query Match 100.0%; Score 29; DB 4; Length 11;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|||||
Db 1 TKPPR 5

RESULT 39

US-09-387-715-52
; Sequence 52, Application US/09387715
; Patent No. 6551574
; GENERAL INFORMATION:
; APPLICANT: Sharma, Shubb
; TITLE OF INVENTION: Tuftsin Metallopeptides Analogs and Uses Thereof
; FILE REFERENCE: 1173/1D794US1
; CURRENT APPLICATION NUMBER: US/09/387,715
; CURRENT FILING DATE: 1999-08-30
; PRIOR APPLICATION NUMBER: PCT/US99/05693
; PRIOR FILING DATE: 1999-03-18
; NUMBER OF SEQ ID NOS: 55
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 52
; LENGTH: 11
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: synthetic
; OTHER INFORMATION: peptide
; NAME/KEY: MOD_RES
; LOCATION: 6
; OTHER INFORMATION: Ahe
; NAME/KEY: modified residue
; LOCATION: 2,4
; OTHER INFORMATION: D-amino acid
US-09-387-715-52

Query Match 100.0%; Score 29; DB 4; Length 11;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
|||||
Db 7 TKPPR 11

RESULT 40

US-08-299-636-39
; Sequence 39, Application US/08299636
; Patent No. 5659041
; GENERAL INFORMATION:
; APPLICANT: POLLAK, Alfred
; APPLICANT: KIRBY, Robert A.
; APPLICANT: DUNN-DUFAULT, Robert
; TITLE OF INVENTION: HYDRAZINO-TYPE RADIONUCLIDE CHELATORS
; TITLE OF INVENTION: HAVING AN NIS CONFIGURATION
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: 3000 K Street, N.W., Suite 500
; CITY: Washington
; STATE: D.C.
; COUNTRY: USA
; ZIP: 20007-5109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/299,636
; FILING DATE: 02-SEP-1994
; CLASSIFICATION: 534
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/092,911
; FILING DATE: 18-JUL-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: BENT, Stephen A.
; REGISTRATION NUMBER: 29,768
; REFERENCE/DOCKET NUMBER: 16777/262/ALLE
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 672-5300
; TELEFAX: (202) 672-5399
; TELEX: 904136
; INFORMATION FOR SEQ ID NO: 39:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 12 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /product= "OTHER"
; OTHER INFORMATION: /note= "The Xaa at position 1 = sulphur"
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 2
; OTHER INFORMATION: /product= "OTHER"
; OTHER INFORMATION: /note= "The Xaa at position 2 = Acn"
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 3
; OTHER INFORMATION: /product= "OTHER"
; OTHER INFORMATION: /note= "The Xaa at position 3 = Mercaptoacetyl"
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 5
; OTHER INFORMATION: /product= "OTHER"

```
; OTHER INFORMATION: /note= "The Xaa at position 5 = nitrogen"
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 6
; OTHER INFORMATION: /product= "OTHER"
; OTHER INFORMATION: /note= "The Xaa at position 6 = methylhydrazino nicotinic acid"
US-08-299-636-39

Query Match      100.0%; Score 29; DB 1; Length 12;
Best Local Similarity 100.0%; Pred. No. 19;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 8 TKPPR 12

RESULT 41
US-09-381-546-13
; Sequence 13, Application US/09381546
; Patent No. 6451976
; GENERAL INFORMATION:
; APPLICANT: Trigen Limited
; TITLE OF INVENTION: BI- OR MULTIFUNCTIONAL MOLECULES BASED ON A DENDROASPIN
; FILE REFERENCE: P41007NO
; CURRENT APPLICATION NUMBER: US/09/381,546
; CURRENT FILING DATE: 1999-09-20
; PRIOR APPLICATION NUMBER: PCT/GB98/00848
; PRIOR FILING DATE: 1998-09-20
; PRIOR APPLICATION NUMBER: GB9705787.1
; PRIOR FILING DATE: 1997-03-20
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 13
; LENGTH: 60
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: modified dendroaspin
US-09-381-546-13

Query Match      100.0%; Score 29; DB 4; Length 60;
Best Local Similarity 100.0%; Pred. No. 82;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 9 TKPPR 13

RESULT 42
US-09-252-991A-25421
; Sequence 25421, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 25421
; LENGTH: 98
; TYPE: PRT
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-25421

; OTHER INFORMATION: /note= "The Xaa at position 5 = nitrogen"
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 6
; OTHER INFORMATION: /product= "OTHER"
; OTHER INFORMATION: /note= "The Xaa at position 6 = methylhydrazino nicotinic acid"
US-08-299-636-39

Query Match      100.0%; Score 29; DB 1; Length 12;
Best Local Similarity 100.0%; Pred. No. 19;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 8 TKPPR 12

RESULT 41
US-09-381-546-13
; Sequence 13, Application US/09381546
; Patent No. 6451976
; GENERAL INFORMATION:
; APPLICANT: Trigen Limited
; TITLE OF INVENTION: BI- OR MULTIFUNCTIONAL MOLECULES BASED ON A DENDROASPIN
; FILE REFERENCE: P41007NO
; CURRENT APPLICATION NUMBER: US/09/381,546
; CURRENT FILING DATE: 1999-09-20
; PRIOR APPLICATION NUMBER: PCT/GB98/00848
; PRIOR FILING DATE: 1998-09-20
; PRIOR APPLICATION NUMBER: GB9705787.1
; PRIOR FILING DATE: 1997-03-20
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 13
; LENGTH: 60
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: modified
; OTHER INFORMATION: dendroaspin
US-09-381-546-13

Query Match      100.0%; Score 29; DB 4; Length 60;
Best Local Similarity 100.0%; Pred. No. 82;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 9 TKPPR 13

RESULT 42
US-09-252-991A-25421
; Sequence 25421, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 25421
; LENGTH: 98
; TYPE: PRT
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-25421

; OTHER INFORMATION: /note= "The Xaa at position 5 = nitrogen"
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 6
; OTHER INFORMATION: /product= "OTHER"
; OTHER INFORMATION: /note= "The Xaa at position 6 = methylhydrazino nicotinic acid"
US-08-299-636-39

Query Match      100.0%; Score 29; DB 4; Length 98;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 72 TKPPR 76

RESULT 43
US-09-252-991A-21156
; Sequence 21156, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 21156
; LENGTH: 136
; TYPE: PRT
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-21156

Query Match      100.0%; Score 29; DB 4; Length 136;
Best Local Similarity 100.0%; Pred. No. 1.7e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 86 TKPPR 90

RESULT 44
US-09-489-039A-8820
; Sequence 8820, Application US/09489039A
; Patent No. 8610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; CURRENT FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 8820
; LENGTH: 420
; TYPE: PRT
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-8820

Query Match      100.0%; Score 29; DB 4; Length 420;
Best Local Similarity 100.0%; Pred. No. 4.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 270 TKPPR 274

RESULT 45
US-09-367-206-1
; Sequence 1, Application US/09367206
```

; Patent No. 6326482
; GENERAL INFORMATION:
; APPLICANT: Genentech, Inc.
; TITLE OF INVENTION: NSP Molecules
; FILE REFERENCE: P1223RIE
; CURRENT APPLICATION NUMBER: US/09/367,206
; CURRENT FILING DATE: 1999-08-09
; PRIOR APPLICATION NUMBER: PCT/US99/08847
; PRIOR FILING DATE: 1999-04-23
; PRIOR APPLICATION NUMBER: US 60/082,767
; PRIOR FILING DATE: 1998-04-23
; PRIOR APPLICATION NUMBER: US 60/113,296
; PRIOR FILING DATE: 1998-12-22
; NUMBER OF SEQ ID NOS: 35
; SEQ ID NO 1
; LENGTH: 576
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-367-206-1

Query Match 100.0%; Score 29; DB 4; Length 576;
Best Local Similarity 100.0%; Pred. No. 6.5e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 211 TKPPR 215

RESULT 46
US-09-367-206-21
; Sequence 21, Application US/09367206
; Patent No. 6326482
; GENERAL INFORMATION:
; APPLICANT: Genentech, Inc.
; TITLE OF INVENTION: NSP Molecules
; CURRENT APPLICATION NUMBER: US/09/367,206
; CURRENT FILING DATE: 1999-08-09
; PRIOR APPLICATION NUMBER: PCT/US99/08847
; PRIOR FILING DATE: 1999-04-23
; PRIOR APPLICATION NUMBER: US 60/082,767
; PRIOR FILING DATE: 1998-04-23
; PRIOR APPLICATION NUMBER: US 60/113,296
; PRIOR FILING DATE: 1998-12-22
; NUMBER OF SEQ ID NOS: 35
; SEQ ID NO 21
; LENGTH: 576
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Mutation of SEQ ID NO:1
US-09-367-206-21

Query Match 100.0%; Score 29; DB 4; Length 576;
Best Local Similarity 100.0%; Pred. No. 6.5e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 211 TKPPR 215

RESULT 47
US-09-367-206-22
; Sequence 22, Application US/09367206
; Patent No. 6326482
; GENERAL INFORMATION:
; APPLICANT: Genentech, Inc.
; TITLE OF INVENTION: NSP Molecules
; FILE REFERENCE: P1223RIE
; CURRENT APPLICATION NUMBER: US/09/367,206
; CURRENT FILING DATE: 1999-08-09

; PRIOR APPLICATION NUMBER: PCT/US99/08847
; PRIOR FILING DATE: 1999-04-23
; PRIOR APPLICATION NUMBER: US 60/082,767
; PRIOR FILING DATE: 1998-04-23
; PRIOR APPLICATION NUMBER: US 60/113,296
; PRIOR FILING DATE: 1998-12-22
; NUMBER OF SEQ ID NOS: 35
; SEQ ID NO 22
; LENGTH: 576
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Mutation of SEQ ID NO:1
US-09-367-206-22

Query Match 100.0%; Score 29; DB 4; Length 576;
Best Local Similarity 100.0%; Pred. No. 6.5e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 211 TKPPR 215

RESULT 48
US-09-367-206-23
; Sequence 23, Application US/09367206
; Patent No. 6326482
; GENERAL INFORMATION:
; APPLICANT: Genentech, Inc.
; TITLE OF INVENTION: NSP Molecules
; FILE REFERENCE: P1223RIE
; CURRENT APPLICATION NUMBER: US/09/367,206
; CURRENT FILING DATE: 1999-08-09
; PRIOR APPLICATION NUMBER: PCT/US99/08847
; PRIOR FILING DATE: 1999-04-23
; PRIOR APPLICATION NUMBER: US 60/082,767
; PRIOR FILING DATE: 1998-04-23
; PRIOR APPLICATION NUMBER: US 60/113,296
; PRIOR FILING DATE: 1998-12-22
; NUMBER OF SEQ ID NOS: 35
; SEQ ID NO 23
; LENGTH: 576
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Mutation of SEQ ID NO:1
US-09-367-206-23

Query Match 100.0%; Score 29; DB 4; Length 576;
Best Local Similarity 100.0%; Pred. No. 6.5e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 211 TKPPR 215

RESULT 49
US-09-857-076-42
; Sequence 42, Application US/08957076C
; Patent No. 6225120
; GENERAL INFORMATION:
; APPLICANT: Kimura, Gary
; APPLICANT: Ruvkun, Gary
; APPLICANT: Patterson, Garth
; APPLICANT: Ogs, Scott
; APPLICANT: Paradis, Suzanne
; APPLICANT: Tissenbaum, Heidi
; APPLICANT: Morris, Jason
; APPLICANT: Kowsek, Allison
; TITLE OF INVENTION: THERAPEUTIC AND DIAGNOSTIC TOOLS FOR
; TITLE OF INVENTION: IMPAIRED GLUCOSE TOLERANCE CONDITIONS

FILE REFERENCE: 00786/351001
CURRENT APPLICATION NUMBER: US/08/857,076C
CURRENT FILING DATE: 1997-05-15
NUMBER OF SEQ ID NOS: 114
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 42
LENGTH: 892
TYPE: PRT
ORGANISM: Caenorhabditis elegans
US-08-857-076-42

Query Match 100.0%; Score 29; DB 3; Length 892;
Best Local Similarity 100.0%; Pred. No. 9.7e+02; Indels 0; Gaps 0;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 TKPPR 5
Db 81 TKPPR 85

RESULT 50
US-08-118-101A-2
Sequence 2, Application US/08118101A
Patent No. 5620892
GENERAL INFORMATION:
APPLICANT: Kurtz, Stephen E.
APPLICANT: Knickerbocker, Aron M.
APPLICANT: McCullough, John R.
TITLE OF INVENTION: A STRAIN OF SACHAROMYCES CEREVISIAE
TITLE OF INVENTION: EXPRESSING THE GENE ENCODING POTASSIUM TRANSPORTER MINK
NUMBER OF SEQUENCES: 16
CORRESPONDENCE ADDRESS:
ADDRESSEE: Burton Rodney
STREET: P.O. Box 4000
CITY: Princeton
STATE: New Jersey
COUNTRY: U.S.A.
ZIP: 08543-4000
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/118,101A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Gaul, Timothy J.
REGISTRATION NUMBER: 33,111
REFERENCE/DOCKET NUMBER: DC27
TELEPHONE: (609) 252-5901
TELEFAX: (609) 252-4526
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1235 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-118-101A-2

Query Match 100.0%; Score 29; DB 1; Length 1235;
Best Local Similarity 100.0%; Pred. No. 1.3e+03; Indels 0; Gaps 0;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 TKPPR 5
Db 438 TKPPR 442

Search completed: March 3, 2004, 12:19:39
Job time : 24 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: March 3, 2004, 12:15:03 ; Search time 39 Seconds
(without alignments)
40.451 Million cell updates/sec

Title: US-09-871-974-2

Perfect score: 29

Sequence: 1 TKPPR 5

Scoring table: BIOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database :

SPTREMBL 25:**

1: sp_archaea:**

2: sp_bacteria:**

3: sp_fungi:**

4: sp_human:**

5: sp_invertebrate:**

6: sp_mammal:**

7: sp_mhc:**

8: sp_organelle:**

9: sp_phage:**

10: sp_plant:**

11: sp_rodent:**

12: sp_virus:**

13: sp_vertebrate:**

14: sp_unclassified:**

15: sp_rvirus:**

16: sp_bacteriaph:**

17: sp_archaeap:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	29	100.0	73	017534	C17534 caenorhabdi
2	29	100.0	106	Q29253	Q29253 sus scrofa
3	29	100.0	115	12 Q98676	Q98676 simian cyto
4	29	100.0	142	16 Q9PG08	Q9PG08 xyiella fas
5	29	100.0	142	16 Q87EU9	Q87EU9 xyiella fas
6	29	100.0	143	13 Q91936	Q91936 acipenser b
7	29	100.0	169	12 Q7T922	Q7T922 human adeno
8	29	100.0	190	15 Q8F4V4	Q8F4V4 leptospira
9	29	100.0	191	10 Q9M250	Q9M250 arabidopsis
10	29	100.0	198	12 Q98674	Q98674 simian cyto
11	29	100.0	212	10 Q8H435	Q8H435 oryza sativ
12	29	100.0	222	16 Q8U6C3	Q8U6C3 agrobacteri
13	29	100.0	226	5 Q45855	Q45855 caenorhabdi
14	29	100.0	227	11 Q810F4	Q810F4 rattus norv
15	29	100.0	227	11 Q7TML1	Q7TML1 mus musculu
16	29	100.0	256	10 Q8SA84	Q8SA84 zea mays (m

17	29	100.0	250	11 Q35455	Q35455 mus musculu
18	29	100.0	278	16 P74248	P74248 synechocyst
19	29	100.0	283	5 Q24859	Q24859 entamoeba h
20	29	100.0	316	4 Q60440	Q60440 homo sapien
21	29	100.0	325	10 Q9FLJ1	Q9FLJ1 arabidopsis
22	29	100.0	326	10 Q9FL70	Q9FL70 arabidopsis
23	29	100.0	334	11 Q8CAP5	Q8CAP5 mus musculu
24	29	100.0	334	16 Q8UKK5	Q8UKK5 agrobacteri
25	29	100.0	351	5 Q23287	Q23287 caenorhabdi
26	29	100.0	354	4 Q60439	Q60439 homo sapien
27	29	100.0	370	16 Q8ZG89	Q8ZG89 yersinia pe
28	29	100.0	375	10 Q80865	Q80865 arabidopsis
29	29	100.0	394	16 Q82NT5	Q82NT5 streptomyce
30	29	100.0	405	16 Q85JB7	Q85JB7 bradyrhizob
31	29	100.0	406	16 Q83A13	Q83A13 coxiella bu
32	29	100.0	409	11 Q9D9R4	Q9D9R4 mus musculu
33	29	100.0	409	11 Q8BHB7	Q8BHB7 mus musculu
34	29	100.0	410	2 Q45945	Q45945 coxiella bu
35	29	100.0	410	2 Q52878	Q52878 coxiella bu
36	29	100.0	415	10 Q9XIF9	Q9XIF9 arabidopsis
37	29	100.0	415	10 Q8S8E1	Q8S8E1 arabidopsis
38	29	100.0	415	17 Q96Z07	Q96Z07 sulfolobus
39	29	100.0	429	10 Q944S3	Q944S3 arabidopsis
40	29	100.0	432	11 Q9D2W5	Q9D2W5 mus musculu
41	29	100.0	442	3 Q06349	Q06349 saccharomyc
42	29	100.0	458	5 Q9BLQ2	Q9BLQ2 meloidogyne
43	29	100.0	460	10 Q94LM4	Q94LM4 oryza sativ
44	29	100.0	491	16 Q828Y7	Q828Y7 streptomyce
45	29	100.0	499	16 Q82GE0	Q82GE0 streptomyce
46	29	100.0	505	16 Q7UT23	Q7UT23 rhodopirell
47	29	100.0	524	11 Q9D318	Q9D318 mus musculu
48	29	100.0	524	11 Q9EQY1	Q9EQY1 rattus norv
49	29	100.0	547	5 Q9VQ69	Q9VQ69 drosophila
50	29	100.0	574	4 Q96HX2	Q96HX2 homo sapien
51	29	100.0	575	5 Q62223	Q62223 caenorhabdi
52	29	100.0	576	4 Q9Y2X4	Q9Y2X4 homo sapien
53	29	100.0	576	4 Q9BRG2	Q9BRG2 homo sapien
54	29	100.0	579	11 Q9EQY2	Q9EQY2 rattus norv
55	29	100.0	585	12 Q7IF18	Q7IF18 rhesus cyto
56	29	100.0	615	11 Q9EQY3	Q9EQY3 rattus norv
57	29	100.0	619	4 Q8NGF6	Q8NGF6 homo sapien
58	29	100.0	669	10 Q8GXJ4	Q8GXJ4 arabidopsis
59	29	100.0	678	16 Q8PKG1	Q8PKG1 xanthomonas
60	29	100.0	722	5 Q9XTS1	Q9XTS1 caenorhabdi
61	29	100.0	776	11 Q8BQ16	Q8BQ16 mus musculu
62	29	100.0	808	2 Q700Z1	Q700Z1 saccharopol
63	29	100.0	809	10 Q8L737	Q8L737 arabidopsis
64	29	100.0	809	10 Q94AB2	Q94AB2 arabidopsis
65	29	100.0	816	10 Q9M2F3	Q9M2F3 arabidopsis
66	29	100.0	832	3 Q871X8	Q871X8 neurospora
67	29	100.0	854	5 Q9U0T9	Q9U0T9 leishmania
68	29	100.0	866	13 Q8JHU7	Q8JHU7 brachydanio
69	29	100.0	892	5 Q95Q17	Q95Q17 caenorhabdi
70	29	100.0	906	4 Q8N4W1	Q8N4W1 homo sapien
71	29	100.0	941	10 Q9SMD9	Q9SMD9 arabidopsis
72	29	100.0	959	10 Q8LGM9	Q8LGM9 arabidopsis
73	29	100.0	962	10 Q23048	Q23048 arabidopsis
74	29	100.0	988	5 Q9N8K2	Q9N8K2 trypanosoma
75	29	100.0	1003	4 Q9HCHO	Q9HCHO homo sapien
76	29	100.0	1003	16 Q98GJ5	Q98GJ5 rhizobium 1
77	29	100.0	1022	5 Q27779	Q27779 schistosoma
78	29	100.0	1047	4 Q60284	Q60284 homo sapien
79	29	100.0	1067	5 Q7YU69	Q7YU69 drosophila
80	29	100.0	1413	5 Q9VK08	Q9VK08 drosophila
81	29	100.0	1430	11 Q8VHK2	Q8VHK2 rattus norv
82	29	100.0	1491	5 Q86BT9	Q86BT9 giardia lam
83	29	100.0	1667	5 Q9VHT6	Q9VHT6 drosophila
84	29	100.0	1706	5 Q9VAF5	Q9VAF5 drosophila
85	26	89.7	33	2 Q7WR15	Q7WR15 escherichia
86	26	89.7	42	16 Q8KE21	Q8KE21 chlorobium
87	26	89.7	47	16 Q9JXF4	Q9JXF4 neisseria m
88	26	89.7	50	16 Q8YTS4	Q8YTS4 anabaena sp
89	26	89.7	65	2 Q7WRH8	Q7WRH8 escherichia

```
90 26 89.7 66 16 Q9JZ50
91 26 89.7 68 16 Q9S279
92 26 89.7 75 16 Q9ZB10
93 26 89.7 81 17 Q8ZWE1
94 26 89.7 86 6 Q9GMP3
95 26 89.7 87 5 Q8XKP6
96 26 89.7 87 16 Q8ZE23
97 26 89.7 88 12 Q8Q405
98 26 89.7 89 10 Q7XNQ7
99 26 89.7 100 17 Q9YCU1
100 26 89.7 101 12 Q9JJA0
101 26 89.7 102 12 Q806X6
102 26 89.7 105 16 Q88KU2
103 26 89.7 110 10 Q8S2F3
104 26 89.7 117 16 Q7UUS8
105 26 89.7 123 2 Q7WX19
106 26 89.7 127 16 Q8DBG4
107 26 89.7 132 12 Q9YMF3
108 26 89.7 139 11 Q80ZS9
109 26 89.7 139 16 Q8KB15
110 26 89.7 140 5 Q9ULJ6
111 26 89.7 140 5 Q8YMP9
112 26 89.7 141 10 Q8S3B2
113 26 89.7 147 10 Q7X6P1
114 26 89.7 148 16 Q9A7H4
115 26 89.7 149 17 Q8TV08
116 26 89.7 153 15 Q06322
117 26 89.7 153 16 P74047
118 26 89.7 154 5 Q9NEH6
119 26 89.7 154 17 Q30100
120 26 89.7 156 16 Q8R8V4
121 26 89.7 157 3 Q87IH3
122 26 89.7 163 5 Q8M2M8
123 26 89.7 167 4 Q8WTX0
124 26 89.7 171 16 Q82N66
125 26 89.7 172 4 Q9NWD7
126 26 89.7 173 12 Q8B184
127 26 89.7 179 16 Q8YXV4
128 26 89.7 183 10 Q7XRQ8
129 26 89.7 183 11 Q9CWA7
130 26 89.7 183 16 Q915Q0
131 26 89.7 185 5 Q61697
132 26 89.7 189 10 Q9ARV1
133 26 89.7 192 12 Q8B8T7
134 26 89.7 192 12 Q7T940
135 26 89.7 192 12 Q7T8D1
136 26 89.7 195 2 Q50902
137 26 89.7 196 16 Q87XW1
138 26 89.7 198 5 Q7YVI3
139 26 89.7 201 16 Q82SQ3
140 26 89.7 213 2 Q7X040
141 26 89.7 213 12 Q8QS79
142 26 89.7 218 2 Q844P8
143 26 89.7 218 2 Q844P7
144 26 89.7 218 2 Q844N6
145 26 89.7 220 12 Q92486
146 26 89.7 220 4 Q9UPE7
147 26 89.7 230 17 Q8ZV71
148 26 89.7 231 10 Q7XVZ1
149 26 89.7 231 12 Q8B6S1
150 26 89.7 232 17 Q9YA94
```

ALIGNMENTS

```
RESULT 1
ID O17534 PRELIMINARY; PRT; 73 AA.
AC O17534; (TrEMBLrel. 05, Created)
DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)
DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
```

```
DE DAP-3 (Fragment).
GN DAP-3.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OC NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=2;
RA Patterson G., Kowek A., Wong A., Liu Y., Ruvkun G.;
RT "The DAF-3 Smad protein antagonizes DAF-7 TGF-beta-receptor signalling
in the C. elegans dauer pathway.";
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF005207; AAB61750.1; -.
FT NON_TER 1 73
FT NON_TER 73 73
SQ SEQUENCE 73 AA; 8061 MW; 7D854125D687F0EE CRC64;
```

Query Match 100.0%; Score 29; DB 5; Length 73;
Best Local Similarity 100.0%; Pred. No. 62;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPR 5
Db 15 TKPR 19

RESULT 2

```
ID Q29253 PRELIMINARY; PRT; 106 AA.
AC Q29253;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE Cytochrome C oxidase polypeptide III (Fragment).
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
OC NCBI_TaxID=9823;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Small intestine;
RX MEDLINE=96327607; PubMed=8672129;
RA Winteroe A.K., Fredholm M., Davies W.;
RT "Evaluation and characterization of a porcine small intestine cDNA
library.";
RL Mamm. Genome 7:509-517(1996).
DR EMBL; F14871; CAA23306.1; -.
FT NON_TER 1 106
FT NON_TER 106 106
SQ SEQUENCE 106 AA; 12552 MW; 4BF91EA407F2EF73 CRC64;
```

Query Match 100.0%; Score 29; DB 6; Length 106;
Best Local Similarity 100.0%; Pred. No. 89;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPR 5
Db 48 TKPR 52

RESULT 3

```
ID Q98676 PRELIMINARY; PRT; 115 AA.
AC Q98676;
DT 01-FEB-1997 (TrEMBLrel. 02, Created)
DT 01-FEB-1997 (TrEMBLrel. 02, Last sequence update)
DT 01-MAR-2002 (TrEMBLrel. 20, Last annotation update)
DE BRLF3 (Fragment).
OS Simian cytomegalovirus.
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Betaherpesvirinae; Cytomegalovirus.
OC NCBI_TaxID=10364;
```

```

RN  SEQUENCE FROM N.A.
RP  PubMed=11725047;
RA  Chang Y., Jeang K., Lietman T., Hayward G.S.;
RT  "Structural organization of the Spliced Immediate-Early Gene Complex
RT  that Encodes the Major Acidic Nuclear (IE1) and Transactivator (IE2)
RT  Proteins of African Green Monkey Cytomegalovirus.";
RL  J. Biomed. Sci. 2:105-130(1995).
RN  [2]
RP  SEQUENCE FROM N.A.
RX  MEDLINE=90080130; PubMed=2152815;
RA  Chang Y.N., Crawford S., Stall J., Rawlins D.R., Jeang K.T.,
RA  Hayward G.S.;
RT  "The palindromic series I repeats in the simian cytomegalovirus major
RT  immediate-early promoter behave as both strong basal enhancers and
RT  cyclic AMP response elements.";
RL  J. Virol. 64:264-277(1990).
RN  [3]
RP  SEQUENCE FROM N.A.
RX  MEDLINE=8719884; PubMed=3033283;
RA  Jeang K.T., Rawlins D.R., Rosenfeld P.J., Shero J.H., Kelly T.J.,
RA  Hayward G.S.;
RT  "Multiple tandemly repeated binding sites for cellular nuclear factor
RT  1 that surround the major immediate-early promoters of simian and
RT  human cytomegalovirus";
RL  J. Virol. 61:1559-1570(1987).
RN  [4]
RP  SEQUENCE FROM N.A.
RX  MEDLINE=93100836; PubMed=8380090;
RA  Chang Y.N., Jeang K.T., Chiou C.J., Chan Y.J., Pizzorno M.,
RA  Hayward G.S.;
RT  "Identification of a large bent DNA domain and binding sites for serum
RT  response factor adjacent to the NF1 repeat cluster and enhancer region
RT  in the major IE94 promoter from simian cytomegalovirus.";
RL  J. Virol. 67:516-529(1993).
DR  EMBL; U38308; A016875.1; -.
FT  NON TER 1
SQ  SEQUENCE 115 AA; 12965 MW; C4251671FBD87AF2 CRC64;

Query Match 100.0%; Score 29; DB 12; Length 115;
Best Local Similarity 100.0%; Pred. No. 97;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 110 TKPPR 114

RESULT 4
Q9PGQ8 PRELIMINARY; PRT; 142 AA.
AC Q9PGQ8;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Hypothetical protein Xf0240.
GN Xf0240.
OS Xylella fastidiosa.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xylella.
OX NCBI_TaxID=2371;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=9a5C;
RX MEDLINE=20365717; PubMed=10910347;
RA Simpson A.J.G., Reinach F.C., Arruda P., Abreu F.A., Acencio M.,
RA Alvaranga R., Alves L.M.C., Araya J.E., Baia G.S., Baptista C.S.,
RA Barros M.H., Bonaccorsi E.D., Bordin S., Bove J.M., Briones M.R.S.,
RA Bueno M.R.P., Camargo A.A., Camargo L.E.A., Carraro D.M., Carrer H.,
RA Colauto N.B., Colombo C., Costa F.F., Costa M.C.R., Costa-Neto C.M.,
RA Coutinho L.L., Cristofani M., Dias-Neto E., Docena C., El-Dorri H.,
RA Facincani A.P., Ferreira A.J.S., Ferreira V.C.A., Ferro J.A.,
RA Fraga J.S., Franco S.C., Franco M.C., Frohme M., Furian L.R.,

```

```

RA Garnier M., Goldman G.H., Goldman M.H.S., Gomes S.L., Gruber A.,
RA Ho P.L., Hoheisel J.D., Junqueira M.L., Kemper E.L., Kitajima J.P.,
RA Krieger J.E., Kuramae E.E., Laigret F., Lambais M.R., Leite L.C.C.,
RA Lemos E.G.M., Lemos M.V.P., Lopes S.A., Lopes C.R., Machado J.A.C.,
RA Machado M.A., Madeira A.M.B.N., Madeira H.M.F., Marino C.L.,
RA Marques M.V., Martins E.A.L., Martins E.M.F., Matsukuma A.Y.,
RA Menck C.F.M., Miracca E.C., Miyaki C.Y., Monteiro-Vitorello C.B.,
RA Moon D.H., Nagai M.A., Nascimento A.L.T.O., Netto L.E.S.,
RA Nhani A. Jr., Nobrega F.G., Nunes L.R., Oliveira M.A.,
RA de Oliveira M.C., de Oliveira R.C., Palmieri D.A., Paris A.,
RA Peixoto B.R., Pereira G.A.G., Pereira H.A. Jr., Pesquero J.B.,
RA Quaggio R.B., Roberto P.G., Rodrigues V. de Rosa A.J.M.,
RA de Rosa V.E. Jr., de Sa R.G., Santelli R.V., Sawasaki H.E.,
RA da Silva A.C.R., da Silva A.M., da Silva F.R., Silva W.A. Jr.,
RA da Silveira J.F., Silvestri M.L.Z., Siqueira W.J., de Souza A.A.,
RA de Souza A.P., Terenzi M.F., Truffi D., Tsai S.M., Tshako M.H.,
RA Vallada H., Van Sluys M.A., Verjovski-Almeida S., Vettore A.L.,
RA Zago M.A., Zatz M., Meidanis J., Setubal J.C.;
RT "The genome sequence of the plant pathogen Xylella fastidiosa.";
RL Nature 406:151-159(2000).
DR EMBL; AE003878; AAF83053.1; -.
DR PIR; C82829; C82829.
DR InterPro; IPR002577; DUF24.
DR Pfam; PF01638; DUF24; 1.
DR ProDom; PD004032; DUF24; 1.
DR Hypothetical protein: Complete proteome.
KW SEQUENCE 142 AA; 15878 MW; CAD3A2B6EDB13E31 CRC64;

Query Match 100.0%; Score 29; DB 16; Length 142;
Best Local Similarity 100.0%; Pred. No. 1.2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 87 TKPPR 91

RESULT 5
Q87EU9 PRELIMINARY; PRT; 142 AA.
AC Q87EU9;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Conserved hypothetical protein.
GN PD0199.
OS Xylella fastidiosa (strain Temecul / ATCC 700964).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Xanthomonadales;
OC Xanthomonadaceae; Xylella.
OX NCBI_TaxID=183190;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22421331; PubMed=12533478;
RA Van Sluys M.A., de Oliveira M.C., Monteiro-Vitorello C.B., Moon D.H.,
RA Miyaki C.Y., Furian L.R., Camargo L.E.A., da Silva A.C.R., da Silva F.R.,
RA Takita M.A., Lemos E.G.M., Machado M.A., Ferro M.I.T., da Silva F.R.,
RA Goldman M.H.S., Goldman G.H., Lemos M.V.F., El-Dorri H., Tsai S.M.,
RA Carrer H., Carraro D.M., de Oliveira R.C., Nunes L.R., Siqueira W.J.,
RA Coutinho L.L., Kimura E.T., Ferro E.S., Harakava R., Kuramae E.E.,
RA Marino C.L., Gigliotti E., Abreu I.L., Alves L.M.C., do Amaral A.M.,
RA Baia G.S., Blanco S.R., Brito M.S., Cannavan F.S., Celestino A.V.,
RA da Cunha A.F., Fenille R.C., Ferro J.A., Formighieri E.F., Kishi L.T.,
RA Leoni S.G., Oliveira A.R., Rosa V.E. Jr., Sasaki F.T., Sena J.A.D.,
RA de Souza A.A., Truffi D., Tsukumo F., Yanai G.M., Zaros L.G.,
RA Civerolo E.L., Simpson A.J.G., Almeida N.F. Jr., Setubal J.C.,
RA Kitajima J.P.;
RT "Comparative analyses of the complete genome sequences of Pierce's
RT disease and citrus variegated chlorosis strains of Xylella
RT fastidiosa"; 195:1018-1026(2003).
RL J. Bacteriol.
DR EMBL; AE012553; AAO28090.1; -.
DR InterPro; IPR002577; DUF24.
DR Pfam; PF01638; DUF24; 1.

```

```
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 142 AA; 15808 MW; D52ED9B736C13E31 CRC64;

Query Match 100.0%; Score 29; DB 16; Length 142;
Best Local Similarity 100.0%; Pred. No. 1.2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
   |||||
Db 87 TKPPR 91

RESULT 6
Q91996 PRELIMINARY; PRT; 143 AA.
AC Q91996;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Thyroid-stimulating hormone precursor.
GN TSH.
OS Acipenser baerii (Siberian sturgeon).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Chondrostei; Acipenseriformes; Acipenseridae;
OC Acipenser.
OX NCBI_TaxID=27689;
   [1]
RN SEQUENCE FROM N.A.
RP TISSUE=Pituitary;
RC MEDLINE=20318422; PubMed=10859263;
RA Querat B., Sellouk A., Salmon C.;
RT "Phylogenetic analysis of the vertebrate glycoprotein hormone family
RT including new sequences of sturgeon (Acipenser baerii) subunits of the
RT two gonadotropins and the thyroid stimulating hormone."
RL Biol. Reprod. 63:222-228(2000).
DR EMBL; AJ251659; CAB93505.1; -.
DR HSSP; P01233; 1HCN.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005179; F:hormone activity; IEA.
DR InterPro; IPR006208; Cys knot.
DR Pfam; PF00007; Cys knot; 1.
DR SMART; SM00068; GHb; 1.
DR PROSITE; PS00261; GLYCO_HORMONE_BETA_1; 1.
DR PROSITE; PS00689; GLYCO_HORMONE_BETA_2; 1.
KW SIGNAL.
FT CHAIN 1 20 POTENTIAL.
FT STRAIN 21 143 THYROID-STIMULATING HORMONE.
SQ SEQUENCE 143 AA; 15566 MW; B79E009F7F0ED315 CRC64;

Query Match 100.0%; Score 29; DB 13; Length 143;
Best Local Similarity 100.0%; Pred. No. 1.2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
   |||||
Db 126 TKPPR 130

RESULT 7
Q7T922 PRELIMINARY; PRT; 169 AA.
AC Q7T922;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Agnoprotein.
OS Human adenovirus B.
OC Viruses; dsDNA viruses, no RNA stage; Adenoviridae; Mastadenovirus.
OX NCBI_TaxID=108098;
   [1]
RN SEQUENCE FROM N.A.
RP STRAIN=35p;
RC
```

```
PubMed=12857895;
RA Vogels R., Zuidgeest D., Van Rijnsoever R., Hartkoorn E., Damen I.,
RA De Bethune M.P., Kostense S., Penders G., Helmus N., Koudstaal W.,
RA Cecchini M., Wetterwald A., Sprangers M., Lemckert A., Ophorst O.,
RA Koel B., Van Meerendonk M., Quax P., Panitti L., Grimbergen J.,
RA Bout A., Goudsmit J., Havenga M.;
RT "Replication-Deficient Human Adenovirus Type 35 Vectors for Gene
RT Transfer and Vaccination: Efficient Human Cell Infection and Bypass of
RT Preexisting Adenovirus Immunity."
RL J. Virol. 77:8263-8271(2003).
DR EMBL; AY271307; AAP92368.1; -.
SQ SEQUENCE 169 AA; 18824 MW; DC18351F66B90119 CRC64;

Query Match 100.0%; Score 29; DB 12; Length 169;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
   |||||
Db 97 TKPPR 101

RESULT 8
Q8F4V4 PRELIMINARY; PRT; 130 AA.
AC Q8F4V4;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Conserved hypothetical protein with UVR motif.
GN LA1935.
OS Leptospira interrogans.
OC Bacteria; Spirochaetes; Spirochaetales; Leptospiraceae; Leptospira.
OX NCBI_TaxID=173;
   [1]
RN SEQUENCE FROM N.A.
RP STRAIN=56601 / Serogroup Icterohaemorrhagiae / Serovar lai;
RA Ren S.;
RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB011366; AAN49134.1; -.
DR GO; GO:0003677; F:DNA binding; IEA.
DR GO; GO:0004518; F:nuclease activity; IEA.
DR GO; GO:0006289; P:nucleotide-excision repair; IEA.
DR InterPro; IPR003729; DUF151.
DR InterPro; IPR001943; UvrB/C.
DR Pfam; PF02577; DUF151; 1.
DR Pfam; PF02151; UVR; 1.
DR PROSITE; PS0151; UVR; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 190 AA; 21142 MW; 9A13B59D646E0AD CRC64;

Query Match 100.0%; Score 29; DB 16; Length 190;
Best Local Similarity 100.0%; Pred. No. 1.6e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
   |||||
Db 52 TKPPR 56

RESULT 9
Q9M250 PRELIMINARY; PRT; 191 AA.
AC Q9M250;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein.
GN F7M19 120.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
```


OX NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Nyakatura G., Partmann B., Dauner D., Sterr W., Holland R.,
 RL Weichselgartner M., Mewes H.W., Rudd S., Lemcke K., Mayer K.F.X.,
 RA Quetier F., Salanoubat M.;
 RL Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA EU Arabidopsis sequencing project;
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AL138643; CAB86482.1; --
 DR PIR; T47369; T47369.
 KW Hypothetical protein.
 SQ SEQUENCE 191 AA; 21741 MW; 26F8764BEECB8A85D CRC64;
 Query Match 100.0%; Score 29; DB 10; Length 191;
 Best Local Similarity 100.0%; Pred. No. 1.6e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 TKPR 5
 Db 78 TKPR 82
 RESULT 10
 ID Q98674 PRELIMINARY; PRT; 198 AA.
 AC Q98674;
 DT 01-FEB-1997 (TREMBLrel. 02, Created)
 DT 01-FEB-1997 (TREMBLrel. 02, Last sequence update)
 DT 01-MAR-2002 (TREMBLrel. 20, Last annotation update)
 DE U08F3.
 OS Simian cytomegalovirus.
 OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
 OC Betaherpesvirinae; Cytomegalovirus.
 OX NCBI_TaxID=10364;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX PubMed=11725047;
 RA Chang Y., Jeang K., Lietman T., Hayward G.S.;
 RT "Structural Organization of the Spliced Immediate-Early Gene Complex
 that Encodes the Major Acidic Nucleic (IE1) and Transactivator (IE2)
 Proteins of African Green Monkey Cytomegalovirus.";
 RL J. Biomed. Sci. 2:105-130 (1995).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90080130; PubMed=2152815;
 RA Chang Y.N., Crawford S., Stall J., Rawlins D.R., Jeang K.T.,
 RA Hayward G.S.;
 RT "The palindromic series I repeats in the simian cytomegalovirus major
 immediate-early promoter behave as both strong basal enhancers and
 cyclic AMP response elements.";
 RL J. Virol. 64:264-277 (1990).
 RN [3]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87198884; PubMed=3033283;
 RA Jeang K.T., Rawlins D.R., Rosenfeld P.J., Shero J.H., Kelly T.J.,
 RA Hayward G.S.;
 RT "Multiple tandemly repeated binding sites for cellular nuclear factor
 1 that surround the major immediate-early promoters of simian and
 human cytomegalovirus.";
 RL J. Virol. 61:1559-1570 (1987).
 RN [4]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93100836; PubMed=9380090;
 RA Chang Y.N., Jeang K.T., Chiou C.J., Chan Y.J., Pizzorno M.,
 RA Hayward G.S.;
 RT "Identification of a large bent DNA domain and binding sites for serum
 response factor adjacent to the NF1 repeat cluster and enhancer region
 in the major IE94 promoter from simian cytomegalovirus.";
 RL J. Virol. 67:516-529 (1993).
 DR EMBL; U38308; AB16873.1; --.

SQ SEQUENCE 198 AA; 22221 MW; DB6CE52D3775B0A CRC64;
 Query Match 100.0%; Score 29; DB 12; Length 198;
 Best Local Similarity 100.0%; Pred. No. 1.6e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 TKPR 5
 Db 193 TKPR 197
 RESULT 11
 ID Q8H435 PRELIMINARY; PRT; 212 AA.
 AC Q8H435;
 DT 01-MAR-2003 (TREMBLrel. 23, Created)
 DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
 DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
 DE P0407H12.35 protein.
 GN P0407H12.35.
 OS Oryza sativa (japonica cultivar-group).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Ehrhartoideae; Oryzoae; Oryza.
 OX NCBI_TaxID=39947;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Nipponbare;
 RA Sasaki T., Matsumoto T., Yamamoto K.;
 RT "Oryza sativa nipponbare (GAS) genomic DNA, chromosome 7, PAC
 clone: P0407H12.35";
 RL Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AP004303; BAC21456.1; --
 SQ SEQUENCE 212 AA; 24492 MW; 0EB7FD1860C63994 CRC64;
 Query Match 100.0%; Score 29; DB 10; Length 212;
 Best Local Similarity 100.0%; Pred. No. 1.8e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 TKPR 5
 Db 108 TKPR 112
 RESULT 12
 ID Q8U6C3 PRELIMINARY; PRT; 222 AA.
 AC Q8U6C3;
 DT 01-JUN-2002 (TREMBLrel. 21, Created)
 DT 01-JUN-2002 (TREMBLrel. 21, Last sequence update)
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
 DE Hypothetical protein Atu4884.
 GN ATU4884 OR AGR_L_19.
 OS Agrobacterium tumefaciens (strain C58 / ATCC 33970).
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
 OC Rhizobiaceae; Rhizobium/Agrobacterium group; Agrobacterium.
 OX NCBI_TaxID=176299;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21608550; PubMed=11743193;
 RA Wood D.W., Setubal J.C., Kaul R., Monks D.E., Kitajima J.P.,
 RA Wood D.W., Setubal J.C., Kaul R., Monks D.E., Almeida N.F. Jr., Woo L.,
 RA Okura Y.K., Zhou Y., Chen L., Wood G.E., Bovee D. Sr.,
 RA Chen Y., Paulsen I.T., Eissen J.A., Karp P.D., Bovee D. Sr.,
 RA Chapman P., Clendinning J., Deatherage G., Gillet W., Grant C.,
 RA Kutayvin T., Levy R., Li M.-J., McClelland E., Palmieri A.,
 RA Raymond C., Rouse G., Saenphimachak C., Wu Z., Romero P., Gordon D.,
 RA Zhang S., Yoo H., Tao Y., Biddle P., Jung M., Krespan W., Perry M.,
 RA Gordon-Kamm B., Liao L., Kim S., Hendrick C., Zhao Z.-Y., Dolan M.,
 RA Chumley P., Tingey S.V., Tomb J.-F., Gordon M.P., Olson M.V.,
 RA Nester E.W.;
 RT "The genome of the natural genetic engineer Agrobacterium tumefaciens
 C58.";
 RL Science 294:2317-2323 (2001).

```

RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=21608551; PubMed=11743124; Miller N., Blanchard M., Mullin L.,
RA Goodner B., Hinkle G., Gattung S., Cao Y., Askenazi M., Halling C., Liu F.,
RA Houlihan K., Gordon J., Vaudin M., Tarchouk O., Epp A., Liu F.,
RA Wollam C., Allinger M., Doughty D., Scott C., Lappas C., Markelz B.,
RA Planagan C., Crowell C., Gursion J., Lomo C., Sear C., Strub G.,
RA Cielo C., Slater S.;
RT "Genome sequence of the plant pathogen and biotechnology agent
RT Agrobacterium tumefaciens C58."
RL Science 254:2323-2328(2001).
DR EMBL; AB009416; AAL45678.1; ALT_INIT.
DR EMBL; AB008198; AKK88561.1; -.
DR PIR; AH3157; AH3157.
DR PIR; G98129; G98129.
DR GO; GO:0003677; F:DNA binding; IEA.
DR GO; GO:0006355; P:regulation of transcription, DNA-dependent; IEA.
DR InterPro; IPR002145; HTH_CoPG.
DR Pfam; PF01402; HTH_4; 1.
RW Hypothetical protein; Complete proteome.
SQ SEQUENCE 222 AA; 24796 MW; C8E8A5C3043B6EF3 CRC64;

Query Match 100.0%; Score 29; DB 16; Length 222;
Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 77 TKPPR 81

RESULT 13
C45855
ID O45855 PRELIMINARY; PRT; 226 AA.
AC O45855;
DT 01-JUN-1998 (TrEMBLrel. 06, Created)
DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE T2E7.1 protein.
DE T2E7.1.
GN Caenorhabditis elegans.
OS Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditidae;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RA Cummings P.N.;
RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=99069613; PubMed=9851916;
RA none;
RT "Genome sequence of the nematode C.elegans: A platform for
RT investigating biology."
RL Science 282:2012-2018(1998).
DR EMBL; Z82284; CAB05287.1; -.
DR PIR; T25360; T25360.
DR WormPep; T2E7.1; CE16505.
SQ SEQUENCE 226 AA; 25159 MW; 1C98B5A2873B6737 CRC64;

Query Match 100.0%; Score 29; DB 5; Length 226;
Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 75 TKPPR 79

RESULT 14
Q810F4
ID Q810F4 PRELIMINARY; PRT; 227 AA.

```

```

AC Q810F4;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE FAM3C-like protein.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
CN NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=Wistar; TISSUE=Bone marrow;
RA Buki K.G., Vaananen K.;
RT "Novel genes in rat bone marrow."
RL Submitted (FEB-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY228475; AAO73558.1; -.
SQ SEQUENCE 227 AA; 24713 MW; 43A84E5B326D44A CRC64;

Query Match 100.0%; Score 29; DB 11; Length 227;
Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 51 TKPPR 55

RESULT 15
Q7TML1
ID Q7TML1 PRELIMINARY; PRT; 227 AA.
AC Q7TML1;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE D6Wsl76e protein.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
CN NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=CZECH II; TISSUE=Breast tumor;
RX MEDLINE=22389257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.W.,
RA Bosak S.A., McSwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
RA Krzywinski M.I., Skalska U., Smailus D.E., Schnerch A., Schein J.E.,
RA Jones S.J., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=CZECH II; TISSUE=Breast tumor;
RA Strausberg R.;
RL Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC055853; AAH55853.1; -.
SQ SEQUENCE 227 AA; 24783 MW; 98F02167EBB68CE1 CRC64;

Query Match 100.0%; Score 29; DB 11; Length 227;
Best Local Similarity 100.0%; Pred. No. 1.9e+02;

```

```
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 51 TKPPR 55

RESULT 16
Q8SAB4 PRELIMINARY; PRT; 256 AA.
AC Q8SAB4;
DT 01-JUN-2002 (TrEMBLrel. 21, Created)
DT 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Z19SD10.20 protein (Fragment).
GS Z19SD10.20.
OS Zea mays (Maize).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC PACAD clade; Panicoideae; Andropogoneae; Zea.
OX NCBI_TaxID=4577;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. B73;
RA Ramakrishna W., Emberton J., SanMiguel P., Bennetzen J.;
RL Submitted (JAN-2002) to the ENBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. B73;
RA Doebley J.;
RL Submitted (JAN-2002) to the ENBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. B73;
RA Liaca V., Linton E.W., Young S., Kovchok S., Messing J.;
RL Submitted (JAN-2002) to the ENBL/GenBank/DBJ databases.
DR EMBL; AF466646; AAL76008.1; -.
DR InterPro; IPR005333; TCP.
DR Pfam; PF03634; TCP; 1.
FT NON TER 256
SQ SEQUENCE 256 AA; 27971 MW; 79596CC55483648E CRC64;

Query Match 100.0%; Score 29; DB 10; Length 256;
Best Local Similarity 100.0%; Pred. No. 2.1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 216 TKPPR 220

RESULT 17
O35455 PRELIMINARY; PRT; 260 AA.
AC O35455;
DT 01-JAN-1998 (TrEMBLrel. 05, Created)
DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Homeobox protein Nkx2.6 (Fragment).
GN NKX2-6 OR NKX2.6.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98145949; PubMed=9486544;
RA Nikolaeva M., Chen X., Lufkin T.;
RT "Nkx2.6 expression is transiently and specifically restricted to the
RT branchial region of pharyngeal-stage mouse embryos.";
RL Mech. Dev. 69:215-218 (1997).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
DR EMBL; AF030113; AAB86406.1; -.
```

```
DR HSSP; P23441; 1FTT.
DR MGD; MGI-97351; Nkx2-6.
DR GO; GO:0005634; C.nucleus; IEA.
DR GO; GO:0003700; F:transcription factor activity; IEA.
DR GO; GO:0006355; P:regulation of transcription, DNA-dependent; IEA.
DR InterPro; IPR001356; Homeobox.
DR Pfam; PF00046; homeobox; 1.
DR PRINTS; PR00024; HOMEBOX.
DR ProDom; PD000010; Homeobox; 1.
DR SMART; SM00389; HOX; 1.
DR PROSITE; PS00027; HOMEBOX_1; 1.
DR PROSITE; PSS0071; HOMEBOX_2; 1.
KW DNA-binding; Homeobox; Nuclear protein.
FT NON TER 1
SQ SEQUENCE 260 AA; 28360 MW; 3264612FA98EBF9B CRC64;

Query Match 100.0%; Score 29; DB 11; Length 260;
Best Local Similarity 100.0%; Pred. No. 2.1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 161 TKPPR 165

RESULT 18
P74248 PRELIMINARY; PRT; 278 AA.
AC P74248;
DT 01-FEB-1997 (TrEMBLrel. 02, Created)
DT 01-FEB-1997 (TrEMBLrel. 02, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein slr1169.
GN SLR1169.
OS Synechocystis sp. (strain PCC 6803).
OC Bacteria; Cyanobacteria; Chroococcales; Synechocystis.
OX NCBI_TaxID=1148;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97061201; PubMed=8905231;
RA Kaneko T., Sato S., Kotani H., Tanaka A., Asamizu E., Nakamura Y.,
RA Miyajima N., Hiroseawa M., Sugita M., Sasamoto S., Kimura T.,
RA Hosouchi T., Matsuno A., Muraki A., Nakazaki N., Naruo K., Okumura S.,
RA Shimo S., Takeuchi C., Wada T., Watanabe A., Yamada M., Yasuda M.,
RA Tabata S.;
RT "Sequence analysis of the genome of the unicellular cyanobacterium
RT Synechocystis sp. strain PCC6803. II. Sequence determination of the
RT entire genome and assignment of potential protein-coding regions.";
RL DNA Res. 3:109-136 (1996).
DR EMBL; D90913; BAA18342.1; -.
DR PIR; S75883; S75883.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 278 AA; 30966 MW; E6503E98DB8C81EA CRC64;

Query Match 100.0%; Score 29; DB 16; Length 278;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 240 TKPPR 244

RESULT 19
Q24859 PRELIMINARY; PRT; 283 AA.
ID Q24859;
AC Q24859;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein (Fragment).
OS Entamoeba histolytica.
OC Eukaryota; Entamoebidae; Entamoeba.
```

```
OX NCBI_TaxID=5759;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=HM-1:IMSS;
RX MEDLINE=95327678; PubMed=7604025;
RA Clark C.G., Roger A.J.;
RT "Direct evidence for secondary loss of mitochondria in Entamoeba
RL histolytica";
RN Proc. Natl. Acad. Sci. U.S.A. 92:6518-6521(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=HM-1:IMSS;
RX Clark C.G.;
RL Submitted (NOV-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: L39933; AAC41578.1; -
DR PIR: T18299; T18299.
KW Hypothetical protein.
FT NON_TER 283
SQ SEQUENCE 283 AA; 32222 MW; DD43E869752F9697 CRC64;

Query Match 100.0%; Score 29; DB 5; Length 283;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 208 TKPPR 212

RESULT 20
O60440 PRELIMINARY; PRT; 316 AA.
AC O60440;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Alpha-actinin-2 associated LIM protein.
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Xia H., Bredt D.S.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
RC -1- SIMILARITY: CONTAINS 1 LIM DOMAIN. THE LIM DOMAIN BINDS 2 ZINC
CC IONS.
CC -1- SIMILARITY: CONTAINS 1 PDZ/DHR DOMAIN.
DR EMBL: AF002282; AAC16672.1; -
DR GO: GO:0007242; P:intracellular signaling cascade; IEA.
DR InterPro: IPR001781; LIM.
DR InterPro: IPR001478; PDZ.
DR InterPro: IPR006643; ZASP.
DR Pfam: PF00412; LIM; 1.
DR Pfam: PF00595; PDZ; 1.
DR ProDom: PD000094; LIM; 1.
DR SMART: SM00132; LIM; 1.
DR SMART: SM00228; PDZ; 1.
DR SMART: SM00735; ZM; 1.
DR PROSITE: PS50023; LIM_DOMAIN_2; 1.
DR PROSITE: PS50106; PDZ; 1.
DR LIM domain; Metal-binding; Zinc.
KW LIM domain; Metal-binding; Zinc.
SQ SEQUENCE 316 AA; 34266 MW; C6D0A5E8A5DA3FF90 CRC64;

Query Match 100.0%; Score 29; DB 4; Length 316;
Best Local Similarity 100.0%; Pred. No. 2.6e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 301 TKPPR 305
```

```
RESULT 21
O9FLJ1 PRELIMINARY; PRT; 325 AA.
AC O9FLJ1;
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Genomic DNA, chromosome 5, P1 clone:MIO24.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; Core eudicots; rosids;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Columbia;
RX MEDLINE=98290546; PubMed=9628582;
RA Sato S., Kaneko T., Kotani H., Nakamura Y., Asamizu E., Miyajima N.,
RA Tabata S.;
RT "Structural analysis of Arabidopsis thaliana chromosome 5. IV.
RT Sequence features of the regions of 1,456,315 bp covered by nineteen
RT physically assigned P1 and TAC clones.";
RL DNA Res. 5:41-54(1998).
DR EMBL: AB010074; BAB11237.1; -
SQ SEQUENCE 325 AA; 35789 MW; 9887D8DF8A81C6AA CRC64;

Query Match 100.0%; Score 29; DB 10; Length 325;
Best Local Similarity 100.0%; Pred. No. 2.7e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
Db 9 TKPPR 13

RESULT 22
O9FL70 PRELIMINARY; PRT; 326 AA.
AC O9FL70;
DT 01-WAR-2001 (TrEMBLrel. 16, Created)
DT 01-WAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Cotton fiber expressed protein 1-like protein.
GN AT5G54300.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; Core eudicots; rosids;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Columbia;
RX MEDLINE=98344145; PubMed=9679202;
RA Kaneko T., Kotani H., Nakamura Y., Sato S., Asamizu E., Miyajima N.,
RA Tabata S.;
RT "Structural analysis of Arabidopsis thaliana chromosome 5. V. Sequence
RT features of the regions of 1,381,565 bp covered by twenty one
RT physically assigned P1 and TAC clones.";
RL DNA Res. 5:131-145(1998).
RN [2]
RP SEQUENCE FROM N.A.
RA Nguyen M., Karlin-Neumann G., Southwick A., Lam B., Miranda M.,
RA Palm C.J., Bowser L., Jones T., Banh J., Carninci P., Chen H.,
RA Cheuk R., Chung M.K., Hayaishizaki Y., Ishida J., Kamiya A., Kawai J.,
RA Kim C., Lin J., Liu S.X., Narusaka M., Pham P.K., Sakano H.,
RA Sakurai T., Satou M., Seki M., Shinn P., Yamada K., Shinozaki K.,
RA Ecker J., Theologis A., Davis R.W.;
RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RA Nguyen M., Karlin-Neumann G., Southwick A., Tripp M., Miranda M.,
RA Palm C.J., Bowser L., Jones T., Banh J., Carninci P., Chen H.,
RA Cheuk R., Chung M.K., Hayaishizaki Y., Ishida J., Kamiya A., Kawai J.,
```

RA Kim C., Lin J., Liu S.X., Narusaka M., Pham P.K., Sakano H.,
 RA Sakurai T., Satou M., Seki M., Shinn P., Yamada K., Shinozaki K.,
 RA Ecker J., Theologis A., Davis R.W.;
 RL Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AB010695; BAB10753.1; -;
 DR EMBL; AY098819; AAM20670.1; -;
 DR EMBL; BT000327; AAN15646.1; -;
 DR InterPro; IPR008480; DUF761.
 DR Pfam; PF05553; DUF761.1;
 SQ SEQUENCE 326 AA; 36448 MW; 6EE00EC2D23B1F5F CRC64;

Query Match 100.0%; Score 29; DB 10; Length 326;
 Best Local Similarity 100.0%; Pred. No. 2.7e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 DB 210 TKPRP 214

RESULT 23
 Q8CAP5 PRELIMINARY; PRT; 334 AA.
 AC Q8CAP5;
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE Hypothetical GRAM domain containing protein.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathia; Muridae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Thymus;
 RX MEDLINE=22354683; PubMed=12466851;
 RA The FANTOM Consortium.
 RA "Analysis of the mouse transcriptome based on functional annotation of
 RT 60,770 full-length cDNAs."
 RL Nature 420:563-573(2002).
 DR EMBL; AK038289; BAC29960.1; -;
 DR InterPro; IPR004182; GRAM_dom.
 DR Pfam; PF02893; GRAM; 1.
 DR SMART; SM00568; GRAM; 1.
 KW Hypothetical protein.
 SQ SEQUENCE 334 AA; 37116 MW; 57AE3045256D81A9 CRC64;

Query Match 100.0%; Score 29; DB 11; Length 334;
 Best Local Similarity 100.0%; Pred. No. 2.7e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 DB 306 TKPRP 310

RESULT 24
 Q8UKK5 PRELIMINARY; PRT; 350 AA.
 AC Q8UKK5;
 DT 01-JUN-2002 (TrEMBLrel. 21, Created)
 DT 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE ATP-dependent DNA ligase.
 GN ATU5097 OR AGR PAT 142.
 OS Agrobacterium tumefaciens (strain C58 / ATCC 33970).
 OC Plasmid AT.
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
 OC Rhizobiaceae; Rhizobium/Agrobacterium group; Agrobacterium.
 OX NCBI_TaxID=176299;
 RN [1]
 RP SEQUENCE FROM N.A.

RA Wood D.W., Setubal J.C., Kaul R., Monks D.E., Kitajima J.P.,
 RA Okura Y.K., Zhou Y., Chen L., Wood G.E., Almeida N.F. Jr., Woo L.,
 RA Chen Y., Paulsen I.T., Eisen J.A., Karp P.D., Hovee D. Sr.,
 RA Chapman P., Clendinning J., Deatherage G., Gillet W., Grant C.,
 RA Kutayavin T., Levy R., Li M.-J., McClelland E., Palmieri A.,
 RA Raymond C., Rouse G., Saenphimmachak C., Wu Z., Romero P., Gordon D.,
 RA Zhang S., Yoo H., Tao Y., Biddle P., Jung M., Krespan W., Perry M.,
 RA Gordon-Kamm B., Liao L., Kim S., Hendrick C., Zhao Z.-Y., Dolan M.,
 RA Chumley F., Tingey S.V., Tomb J.-F., Gordon M.P., Olson M.V.,
 RA Nester E.W.;
 RA "The genome of the natural genetic engineer Agrobacterium tumefaciens
 RT C58.";
 RL Science 294:2317-2323(2001).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21608551; PubMed=11743194;
 RA Goodner B., Hinkle G., Gattung S., Miller N., Blanchard M., Mullin L.,
 RA Qurollo B., Goldman B.S., Cao Y., Askenazi M., Halling C., Mullin L.,
 RA Houmlel K., Gordon J., Vaudin M., Tatchouk O., Epp A., Liu F.,
 RA Wollam C., Allinger M., Doughty D., Scott C., Lappas C., Markelz B.,
 RA Flanagan C., Crowell C., Gurson J., Lomo C., Sear C., Strub G.,
 RA Cielo C., Slater S.;
 RA "Genome sequence of the plant pathogen and biotechnology agent
 RT Agrobacterium tumefaciens C58.";
 RL Science 294:2323-2328(2001).
 DR EMBL; AE008934; AAL45787.1; -;
 DR EMBL; AE007882; AAK90473.1; -;
 DR PIR; AE3171; AE3171.
 DR GO; GO:0046821; C:extrachromosomal DNA; IEA.
 DR GO; GO:0005524; F:ATP binding; IEA.
 DR GO; GO:0003910; F:DNA ligase (ATP) activity; IEA.
 DR GO; GO:0016874; F:ligase activity; IEA.
 DR GO; GO:0006310; P:DNA recombination; IEA.
 DR GO; GO:0006281; P:DNA repair; IEA.
 DR GO; GO:0006260; P:DNA replication; IEA.
 DR InterPro; IPR000977; DNA_ligase.
 DR Pfam; PF01068; DNA_ligase; 1.
 DR Pfam; PF04679; DNA_ligase_A_C; 1.
 DR PROSITE; PS00160; DNA_LIGASE_A3; 1.
 KW Ligase; Plasmid; Complete proteome.
 SQ SEQUENCE 350 AA; 39363 MW; 32C15E7081D8233E CRC64;

Query Match 100.0%; Score 29; DB 16; Length 350;
 Best Local Similarity 100.0%; Pred. No. 2.9e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 DB 2 TKPRP 6

RESULT 25
 Q23287 PRELIMINARY; PRT; 351 AA.
 AC Q23287;
 DT 01-NOV-1996 (TrEMBLrel. 01, Created)
 DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Hypothetical protein (PIP-1).
 GN ZC404.8 OR PIP-1.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
 OC Rhabditidae; Peloderinae; Caenorhabditis.
 OX NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Bristol N2;
 RX MEDLINE=99089613; PubMed=9851916;
 RA None;
 RA "Genome sequence of the nematode C. elegans: a platform for
 RT investigating biology. The C. elegans Sequencing Consortium.";
 RL Science 282:2012-2018(1998).

RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Bristol N2;
 RA Bentley D., Le T.T.;
 RT "The sequence of C. elegans cosmid ZC404.";
 RL Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Bristol N2;
 RA Waterston R.;
 RT "Direct Submission.";
 RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RA Ogura K., Kishimoto N., Mitani S., Gengyo-Ando K., Kohara Y.;
 RT "Translational control of maternal glp-1 mRNA by POS-1 and its
 interacting protein SPN-4 in Caenorhabditis elegans.";
 RL Development 0:0-0(2003).
 DR EMBL; U53363; AAA97963.1; -.
 DR EMBL; AB052819; BAC65239.1; -.
 DR PIR; T29369; T29369.
 DR HSSP; P09651; 1HAI.
 DR WormPep; ZC404.8; CE07558.
 DR GO; GO:0003676; F:nuclieic acid binding; IEA.
 DR InterPro; IPR000504; RNA_rec_mot.
 DR Pfam; PF00076; xzm; 1.
 DR SMART; SM00360; RRM; 1.
 DR PROSITE; PS0102; RRM; 1.
 DR PROSITE; PS00030; RRM_RNP_1; 1.
 KW Hypothetical protein.
 SQ SEQUENCE 351 AA; 39213 MW; A83B7159472951E CRC64;
 Query Match 100.0%; Score 29; DB 5; Length 351;
 Best Local Similarity 100.0%; Pred. No. 2.9e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 199 TKPPR 203

DR PROSITE; PS50106; PDZ; 1.
 KW LIM domain; Metal-binding; Zinc.
 SQ SEQUENCE 364 AA; 39231 MW; EAAA2A75466D3E12 CRC64;
 Query Match 100.0%; Score 29; DB 4; Length 364;
 Best Local Similarity 100.0%; Pred. No. 3e+02; 0; Indels 0; Gaps 0;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 349 TKPPR 353

RESULT 27
 Q8ZG89 PRELIMINARY; PRT; 370 AA.
 ID Q8ZG89
 AC Q8ZG89
 DT 01-MAR-2002 (TrEMBLrel. 20, Created)
 DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Putative iron-sulfur binding protein (Hypothetical protein).
 GN YP01417 OR Y2752.
 OS Yersinia pestis.
 CC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
 OC Enterobacteriaceae; Yersinia.
 OX NCBI_TaxID=632;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CO-92 / Biovar Orientalis;
 RX MEDLINE=21470413; PubMed=11586360;
 RA Parkhill J., Wren B.W., Thomson N.R., Titball R.W., Holden M.T.G.,
 RA Prentice M.B., Sebahia M., James K.D., Brooks K., Cerdano-Tarraga A.M.,
 RA Baker S., Baeham D., Bentley S.D., Davies R.M., Davis P., Dougan G.,
 RA Chillingworth T., Cronin A., Cronin A., Davies R.M., Dougan G.,
 RA Felwell T., Hamlin N., Holroyd S., Jagels K., Karlyshev A.V.,
 RA Leather S., Moule S., Oyston P.C.F., Quail M., Rutherford K.,
 RA Simmonds M., Skelton J., Stevens K., Whitehead S., Barrell B.G.;
 RT "Genome sequence of Yersinia pestis, the causative agent of plague.";
 RL Nature 413:523-527(2001).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=KIMS / Biovar Mediaevalis;
 RX MEDLINE=22137863; PubMed=12142430;
 RA Deng W., Burland V., Plunkett G. III, Boutin A., Mayhew G.F., Liss P.,
 RA Perna N.T., Rose D.J., Mau B., Zhou S., Schwartz D.C.,
 RA Fetherston J.D., Lindler L.E., Brubaker R.R., Plano G.V.,
 RA Straley S.C., McDonough K.A., Nilles M.L., Matson J.S., Blattner P.R.,
 RA Perry R.D.;
 RT "Genome sequence of Yersinia pestis KIM.";
 RL J. Bacteriol. 184:4601-4611(2002).
 DR EMBL; AJ414148; CAC90246.1; -.
 DR EMBL; AE013878; RAM86304.1; -.
 DR PIR; AC0173; AC0173.
 DR GO; GO:0005489; F:electron transporter activity; IEA.
 DR GO; GO:0006118; P:electron transport; IEA.
 DR InterPro; IPR006058; 2Fe2S fd BS.
 DR InterPro; IPR001041; Ferredoxin.
 DR InterPro; IPR005302; MOSC.
 DR InterPro; IPR005303; MOSC_N.
 DR Pfam; PF00111; fer2; 1.
 DR Pfam; PF03473; MOSC; 1.
 DR Pfam; PF03476; MOSC_N; 1.
 DR PROSITE; PS00197; 2Fe2S FERREDOXIN; 1.
 KW Hypothetical protein; Complete proteome.
 SQ SEQUENCE 370 AA; 40747 MW; CF6246C4DC713138 CRC64;

Query Match 100.0%; Score 29; DB 16; Length 370;
 Best Local Similarity 100.0%; Pred. No. 3e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 266 TKPPR 270

```

RESULT 29
ID Q80865 PRELIMINARY; PRT; 375 AA.
AC Q80865;
DT 01-NOV-1998 (TrEMBLrel. 08, Created)
DT 01-NOV-1998 (TrEMBLrel. 08, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein.
GN ATG31010.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;
RX MEDLINE=20083487; PubMed=10617197;
RA Lin X.; Kaul S., Rounsley S.D., Shea T.P., Benito M.-I., Town C.D.,
RA Fujii C.Y., Mason T.M., Bowman C.L., Barnstead M.E., Feldblum T.V.,
RA Buell C.R., Ketchum K.A., Lee J.J., Rensing C.M., Koo H., Moffat K.S.,
RA Cronin L.A., Shen M., Vanaken S.B., Umayam L., Tallon L.J., Gill J.B.,
RA Adams M.D., Carrera A.J., Creasy T.H., Goodman H.M., Somerville C.R.,
RA Coppenhaver G.P., Preuss D., Nierman W.C., White O., Eisen J.A.,
RA Salzberg S.L., Fraser C.M., Venter J.C.;
RA "Sequence and analysis of chromosome 2 of the plant Arabidopsis
thaliana";
RT Nature 402:761-768(1999).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;
RX MEDLINE=20083487; PubMed=10617197;
RA Lin X.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
CC -!- SIMILARITY: BELONGS TO THE SER/THR FAMILY OF PROTEIN KINASES.
DR EMBL; AC004669; AAC20735.1; -.
DR PIR; D84715; D84715.
DR HSP; P12931; 1FWK.
DR GO; GO:0005524; F:ATP binding; IEA.
DR GO; GO:0004674; F:protein serine/threonine kinase activity; IEA.
DR GO; GO:0004713; F:protein-tyrosine kinase activity; IEA.
DR GO; GO:0016740; F:transferase activity; IEA.
DR GO; GO:0006468; P:protein amino acid phosphorylation; IEA.
DR InterPro; IPR000719; Prot_kinase.
DR InterPro; IPR008271; Ser_thr_kinase.
DR Pfam; PF00069; Pkinase; 1.
DR PRINTS; PR00109; TYRKINASE.
DR PRODOM; PD000001; Prot_kinase; 1.
DR PROSITE; PS00107; PROTEIN_KINASE_ATP; 1.
DR PROSITE; PS00111; PROTEIN_KINASE_DOM; 1.
DR PROSITE; PS0108; PROTEIN_KINASE_ST; 1.
KW Hypothetical protein; ATP-binding; Kinase;
KW Serine/threonine-protein kinase; Transferase.
SQ SEQUENCE 375 AA; 42588 MW; C1F40CA3A3DE77C5 CRC64;

Query Match 100.0%; Score 29; DB 10; Length 375;
Best Local Similarity 100.0%; Pred. No. 3.1e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 121 TKPPR 125

RESULT 29
ID Q82NT5 PRELIMINARY; PRT; 394 AA.
AC Q82NT5;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein.
GN Savi206.
OS Streptomyces avermitilis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycineae; Streptomycetaceae; Streptomyces.
OX NCBI_TaxID=33903;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
RX MEDLINE=21477403; PubMed=11572948;
RA Omura S., Ikeda H., Ishikawa J., Hanamoto A., Takahashi C.,
RA Shinose M., Shiba T., Sakaki Y., Hattori M.,
RA Kikuchi H., Shiba T., Sakaki Y., Hattori M.;
RT "Genome sequence of an industrial microorganism Streptomyces
avermitilis: deducing the ability of producing secondary
metabolites";
RT Proc. Natl. Acad. Sci. U.S.A. 98:12215-12220(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=MA-4680 / ATCC 31267 / NCIMB 12804 / NRRL 8165;
RX MEDLINE=22608306; PubMed=12692562;
RA Ikeda H., Ishikawa J., Hanamoto A., Shinose M., Kikuchi H., Shiba T.,
RA Sakaki Y., Hattori M., Omura S.;
RT "Complete genome sequence and comparative analysis of the industrial
microorganism Streptomyces avermitilis";
RT Nat. Biotechnol. 21:526-531(2003).
DR EMBL; AP005025; BAC68916.1; -.
DR GO; GO:0016491; F:oxidoreductase activity; IEA.
DR GO; GO:0006118; P:electron transport; IEA.
DR InterPro; IPR006030; Acyl-CoA_dh; 1.
DR Pfam; PF00441; Acyl-CoA_dh; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 394 AA; 42238 MW; B8CF41D0C94F29D2 CRC64;

Query Match 100.0%; Score 29; DB 16; Length 394;
Best Local Similarity 100.0%; Pred. No. 3.2e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 386 TKPPR 390

RESULT 30
ID Q89JB7 PRELIMINARY; PRT; 405 AA.
AC Q89JB7;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE BIL5366 protein.
GN BIL5366
OS Bradyrhizobium japonicum.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Bradyrhizobiaceae; Bradyrhizobium.
OX NCBI_TaxID=375;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=USDA 110;
RX MEDLINE=22484998; PubMed=12597275;
RA Kaneko T., Nakamura Y., Sato S., Minamisawa K., Uchiyama T.,
RA Sasamoto S., Watanabe A., Idesawa K., Iriguchi M., Kawashima K.,
RA Kohara M., Matsumoto M., Shimpo S., Tsuruoka H., Wada T., Yamada M.,
RA Tabata S.;
RT "Complete genomic sequence of nitrogen-fixing symbiotic bacterium
Bradyrhizobium japonicum USDA110";
RT DNA Res. 9:189-197(2002).
DR EMBL; AP005954; BAC50631.1; -.
DR GO; GO:0004730; F:pseudouridylyate synthase activity; IEA.
DR GO; GO:0003723; F:RNA binding; IEA.
DR InterPro; IPR006145; Pseudou synth.
DR InterPro; IPR006224; Rlu_synth.
DR InterPro; IPR002942; S4.

```

```
DR Pfam; PF00849; PseudoU_synth_2; 1.
DR Pfam; PF01479; S4; 1.
DR PROSITE; PS01129; PSI_RLU; 1.
DR PROSITE; PS00889; S4; 1.
DR KW Complete proteome.
SQ SEQUENCE 405 AA; 45209 MW; 2BEE412B47BE36F5 CRC64;

Query Match 100.0%; Score 29; DB 16; Length 405;
Best Local Similarity 100.0%; Pred. No. 3.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 42 TKPPR 46

RESULT 31
Q83A13 PRELIMINARY; PRT; 406 AA.
AC Q83A13;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Site specific recombinase, phage integrase family.
GN CBUA0010.
OS Cxiella burnetii.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Legionellales;
OC Cxiellaceae; Cxiella.
OC NCBI_TaxID=777;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=Nine Mile Phase I / RSA 493;
RX MEDLINE=22608657; PubMed=12704232;
RA Seshadri R., Paulsen I.T., Eisen J.A., Read T.D., Nelson K.E.,
RA Nelson W.C., Ward N.L., Tettelin H., Daviden T.M., Beanan M.J.,
RA DeBoy R.T., Daugherty S.C., Brinkac L.M., Madupu R., Dodson R.J.,
RA Khouri H.M., Lee K.H., Carty H.A., Scanlan D., Heinzen R.A.,
RA Thompson H.A., Samuel J.E., Fraser C.M., Scallan D., Heidelberg J.F.;
RA "Complete genome sequence of the Q-fever pathogen, Cxiella
RT burnetii."
RL Proc. Natl. Acad. Sci. U.S.A. 100:5455-5460 (2003).
DR EMBL; A3016829; AAO91587.1; -.
DR TIGR; CBUA0010; -.
DR GO; GO:0046821; C:extrachromosomal DNA; IEA.
DR GO; GO:0003677; F:DNA binding; IEA.
DR GO; GO:0015074; P:DNA integration; IEA.
DR GO; GO:0006310; P:DNA recombination; IEA.
DR InterPro; IPR002104; Phage integrase.
DR Pfam; PF00589; Phage integrase; 1.
DR KW Plasmid; Complete proteome.
SQ SEQUENCE 406 AA; 47399 MW; 1D40014C4500A23E CRC64;

Query Match 100.0%; Score 29; DB 16; Length 406;
Best Local Similarity 100.0%; Pred. No. 3.3e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 98 TKPPR 102

RESULT 32
Q9D9R4 PRELIMINARY; PRT; 409 AA.
AC Q9D9R4;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
DE 1700030J22Rik protein.
GN 1700030J22Rik.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
```



```
Db 358 TKPPR 362

RESULT 34
ID Q45945 PRELIMINARY; PRT; 410 AA.
AC Q45945; Q45901;
DT 01-NOV-1998 (TRENBLrel. 01, Created)
DT 01-NOV-1998 (TRENBLrel. 01, Last sequence update)
DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)
DE CRF 410.
OS Coccidia burnetii.
OG Plasmid QPH1.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Legionellales;
OC Coccidiaceae; Coccidia.
OX NCBI_TaxID=777;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=NINE MILE PHASE I;
RX MEDLINE=95145653; PubMed=7843345;
RA Thiele D., Willems H., Haas M., Krauss H.;
RT "Analysis of the entire nucleotide sequence of the cryptic plasmid
RL Qph1 from Coccidia burnetii.";
RL Eur. J. Epidemiol. 10:413-420(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=NINE MILE PHASE I;
RA Thiele D.;
RL Submitted (OCT-1993) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=SCURRY Q217;
RA Rittner M., Thiele D., Willems H.;
RT Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
RL EMBL; X75356; CAA53126.1; -.
DR EMBL; X93204; CAA63678.1; -.
DR PIR; S38238; S38238.
DR GO; GO:0046821; C:extrachromosomal DNA; IEA.
DR GO; GO:0003677; F:DNA binding; IEA.
DR GO; GO:0015074; P:DNA recombination; IEA.
DR GO; GO:0006310; P:DNA integration; IEA.
DR InterPro; IPR002104; Phage integrase.
DR Pfam; PF00589; Phage integrase; 1.
KW Plasmid.
SQ SEQUENCE 410 AA; 47829 MW; 9A48C3714B330C2 CRC64;

Query Match 100.0%; Score 29; DB 2; Length 410;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 102 TKPPR 106

RESULT 35
ID Q52878 PRELIMINARY; PRT; 410 AA.
AC Q52878;
DT 01-JUN-1998 (TRENBLrel. 06, Created)
DT 01-JUN-1998 (TRENBLrel. 06, Last sequence update)
DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)
DE Hypothetical protein.
OS Coccidia burnetii.
OG Plasmid QPH1, and Plasmid QpDV.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Legionellales;
OC Coccidiaceae; Coccidia.
OX NCBI_TaxID=777;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PRISCILLA Q177;
RA Lautenschlaeger S., Jaeger C., Willems H., Baljer G.;
RT Submitted (MAR-1998) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=Ril40;
RA Radomski K.J., Willems H., Lautenschlaeger S., Jaeger C., Baljer G.;
RT "Sequence of QpDV plasmid.";
RL Submitted (FEB-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; Y15898; CAA75839.1; -.
DR EMBL; AF131076; AAD33493.1; -.
DR GO; GO:0046821; C:extrachromosomal DNA; IEA.
DR GO; GO:0003677; F:DNA binding; IEA.
DR GO; GO:0015074; P:DNA integration; IEA.
DR GO; GO:0006310; P:DNA recombination; IEA.
DR InterPro; IPR002104; Phage integrase.
DR Pfam; PF00589; Phage integrase; 1.
KW Hypothetical protein; Plasmid.
SQ SEQUENCE 410 AA; 47828 MW; 7A49E15719D330CA CRC64;

Query Match 100.0%; Score 29; DB 2; Length 410;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 102 TKPPR 106

RESULT 36
ID Q9XIF9 PRELIMINARY; PRT; 415 AA.
AC Q9XIF9;
DT 01-NOV-1999 (TRENBLrel. 12, Created)
DT 01-NOV-1999 (TRENBLrel. 12, Last sequence update)
DT 01-JUN-2003 (TRENBLrel. 24, Last annotation update)
DE T10P12.9 protein.
GN T10P12.9.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;
RA Federspiel N.A., Palm C.J., Conway A.B., Conn L., Hansen N.F.,
RA Alafafi H., Araujo R., Huizar L., Rowley D., Buehler E., Dunn P.,
RA Gonzalez A., Kremenetskaia I., Kim C., Lenz C., Li J., Liu S.,
RA Luros S., Schwartz J., Shinn P., Toriumi M., Vysotskaya V.S.,
RA Walker M., Yu G., Ecker J., Theologis A., Davis R.W.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC007203; AAD39275.1; -.
DR PIR; F96499; F96499.
DR HSSP; P50586; LC8Z.
DR InterPro; IPR000007; Tubby.
DR Pfam; PF01167; Tub; 1.
DR PRINTS; PR01573; SUPERTUBBY.
DR PROSITE; PS01200; TUB_1; 1.
SQ SEQUENCE 415 AA; 46492 MW; 38774871B95D8770 CRC64;

Query Match 100.0%; Score 29; DB 10; Length 415;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 309 TKPPR 313

RESULT 37
ID Q8S8E1 PRELIMINARY; PRT; 415 AA.
AC Q8S8E1;
DT 01-JUN-2002 (TRENBLrel. 21, Created)
DT 01-JUN-2002 (TRENBLrel. 21, Last sequence update)
```

Qy 1 TKPPR 5

RESULT 40

Q9D2W5 PRELIMINARY; PRT; 432 AA.
 AC Q9D2W5;
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
 DT 01-NOV-2003 (TrEMBLrel. 23, Last annotation update)
 DE 9130427A09Rik protein.
 DE 9130427A09Rik.
 GN Mus musculus (Mouse).
 OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cecum;
 RX MEDLINE=21085660; PubMed=11217851;
 RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
 RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
 RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
 RA Saito T., Okazaki Y., Gajobori T., Bono H., Kasukawa T., Saito R.,
 RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
 RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
 RA Kuehl P., Lewis S., Mateu Y., Nikaigo I., Pesole G., Quackenbush J.,
 RA Schriml L.M., Staubli F., Suzuki R., Tonita M., Wagner L., Washio T.,
 RA Sakai K., Okido T., Furuno M., Aono H., Baidarelli R., Baren G.,
 RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
 RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gairola M.,
 RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
 RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
 RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
 RA Sasaki H., Sato K., Schenbach C., Seta T., Shibata Y., Storch K.-F.,
 RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
 RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Khtsuki S.,
 RA Hayashizaki Y.;
 RT "Functional annotation of a full-length mouse cDNA collection."
 RL Nature 409:685-690(2001).
 DR EMBL; AK018691; BAB31347.1; -;
 DR MGD; MGI:1914815; 9130427A09Rik.
 DR InterPro; IPR004182; GRAM_dom.
 DR Pfam; PF02893; GRAM; 1.
 DR SMART; SM00568; GRAM; 1.
 SQ SEQUENCE 432 AA; 47932 MW; 2D9898519F2B544B CRC64;

Query Match 100.0%; Score 29; DB 11; Length 432;
 Best Local Similarity 100.0%; Pred. No. 3.5e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 Db 300 TKPRP 304

RESULT 41

Q06349 PRELIMINARY; PRT; 442 AA.
 AC Q06349;
 DT 01-NOV-1996 (TrEMBLrel. 01, Created)
 DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE Chromosome IV COSMID 9481.
 GN YDR370C OR D9481.14.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
 OX NCBI_TaxID=4932;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C;
 RA Ding H.;
 RL Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.

Query Match 100.0%; Score 29; DB 5; Length 458;
 Best Local Similarity 100.0%; Pred. No. 3.7e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 Db 168 TKPRP 172

RESULT 43

Q94LM4

RC STRAIN=S288C;
 RA Johnston M., Andrews S., Brinkman R., Cooper J., Ding H., Du Z.,
 RA Favell A., Fulton L., Gattung S., Greco T., Kirsten J., Kucaba T.,
 RA Hallsworth K., Hawkins J., Hillier L., Jier M., Johnson D.,
 RA Johnston L., Langston Y., Latreille P., Le T., Mardis E., Meneses S.,
 RA Miller N., Nhan M., Pauley A., Peluso D., Rifken L., Riles L.,
 RA Taich A., Trevasxis E., Vignati D., Wilcox L., Wohldman P., Vaudin M.,
 RA Wilson R., Waterston R.;
 RL Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C;
 RA Waterston R.;
 RL Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C;
 RA Jia Y., Cherry J.M.;
 RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
 DR EMBL; U28373; AAB64806.1; -;
 DR PIR; S61165; S61165.
 DR SGD; S0002778; YDR370C.
 SQ SEQUENCE 442 AA; 50492 MW; 504F7C6AA40B50B4 CRC64;

Query Match 100.0%; Score 29; DB 3; Length 442;
 Best Local Similarity 100.0%; Pred. No. 3.6e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 Db 81 TKPRP 85

RESULT 42

Q9BLQ2 PRELIMINARY; PRT; 458 AA.
 AC Q9BLQ2;
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
 DT 01-OCT-2001 (TrEMBLrel. 18, Last annotation update)
 DE Putative avirulence protein precursor.
 GN MAP-1.
 OS Meloidogyne incognita (southern root-knot nematode).
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Tylenchida; Tylenchina;
 OC Tylenchoidea; Heteroderidae; Meloidogyninae; Meloidogyne.
 OX NCBI_TaxID=6306;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21034633; PubMed=1194874;
 RA Semblat J.P., Rosso M.N., Hussey R.S., Abad P., Castagnone-Sereno P.;
 RT "Molecular cloning of a cDNA encoding an amphid-secreted putative
 RT avirulence protein from the root-knot nematode Meloidogyne
 RT incognita";
 RT Mol. Plant Microbe Interact. 14:72-79(2001).
 DR EMBL; AJ278663; CAC27774.1; -;
 DR InterPro; IPR002965; P-rich_extensn.
 DR PRINTS; PR01217; PRICHEXTENS.
 KW SIGNAL.
 FT SIGNAL 1 17 POTENTIAL.
 FT CHAIN 18 458 PUTATIVE AVIRULENCE PROTEIN.
 SQ SEQUENCE 458 AA; 48951 MW; 982AD87D0818815F CRC64;

Query Match 100.0%; Score 29; DB 5; Length 458;
 Best Local Similarity 100.0%; Pred. No. 3.7e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPRP 5
 Db 168 TKPRP 172

RESULT 46

Q7UT23

ID Q7UT23 PRELIMINARY; PRT; 505 AA.
 AC Q7UT23;
 DT 01-OCT-2003 (TrEMBLrel. 25, Created)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE 4-alpha-glucanotransferase (EC 2.4.1.25).
 GN MALQ OR RB4161.
 OS Rhodospirillum rubrum.
 OC Bacteria; Planctomycetes; Planctomycetacia; Planctomycetales;
 OC Planctomycetaceae; Pirellula.
 OX NCBI_TaxID=117;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=1;
 RX MEDLINE=22735913; PubMed=12835416;
 RX Glocner F.O., Kube M., Bauer M., Teeling H., Lombardot T.,
 RA Ludwig W., Gade D., Beck A., Borzym K., Heitmann K., Rabus R.,
 RA Schlesner H., Amann R., Reinhardt R.,
 RT "Complete genome sequence of the marine planctomycete Pirellula sp.
 strain 1.";
 RL Proc. Natl. Acad. Sci. U.S.A. 100:8298-8303(2003).
 RL EMBL; BX294140; CAD73618.1; -.
 KW Glycosyltransferase; Transferase; Complete proteome.
 SQ SEQUENCE 505 AA; 57353 MW; 5470834CCAF15RFF CRC64;

Query Match 100.0%; Score 29; DB 16; Length 505;
 Best Local Similarity 100.0%; Pred. No. 4.1e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

QY 1 TKPPR 5
 Db 121 TKPPR 125

RESULT 47

Q9D318 PRELIMINARY; PRT; 514 AA.
 AC Q9D318;
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
 DT 01-WAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE 9030613F08Rik protein.
 GN 9030613F08Rik.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Colon;
 RX MEDLINE=21085660; PubMed=11217851;
 RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
 RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
 RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
 RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
 RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
 RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
 RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
 RA Schriml L.M., Stauber F., Suzuki R., Tomita M., Wagner L., Washio T.,
 RA Saki K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
 RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
 RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
 RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
 RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
 RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
 RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
 RA Suzuki H., Toyokawa K., Wang K.H., Weitz C., Whitaker C., Wilming L.,
 RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohetsuki S.,
 RA Hayashizaki Y.;
 RT "Functional annotation of a full-length mouse cDNA collection.";
 RL Nature 409:685-690(2001).
 DR EMBL; AK018551; BAB31270.1; -.

DR MGD; MGI:1921790; 9030613F08Rik.
 DR InterPro; IPR004182; GRAM_dom.
 DR Pfam; PF02893; GRAM; 1.
 DR SMART; SM00568; GRAM; 1.
 SQ SEQUENCE 514 AA; 56478 MW; CSA5042F9D228B84 CRC64;

Query Match 100.0%; Score 29; DB 11; Length 514;
 Best Local Similarity 100.0%; Pred. No. 4.2e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 Db 382 TKPPR 386

RESULT 48

Q9EQY1 PRELIMINARY; PRT; 524 AA.
 AC Q9EQY1;
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)
 DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE Transcription factor Elf-1.
 GN ELF-1.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21077473; PubMed=11210123;
 RA Nishiyama C., Takahashi K., Nishiyama M., Okumura K., Ra C.,
 RA Ohtake Y., Yokota T.;
 RT "Polymorphism of transcription factor Elf-1 affecting its regulatory
 function in transcription.";
 RL Biosci. Biotechnol. Biochem. 64:2601-2607(2000).
 DR EMBL; AB030217; BAB20035.1; -.
 DR HSSP; P28324; IBC8.
 DR GO; GO:0005634; C:nucleus; IEA.
 DR GO; GO:0003700; F:transcription factor activity; IEA.
 DR GO; GO:0006355; P:regulation of transcription, DNA-dependent; IEA.
 DR InterPro; IPR000418; Ets.
 DR InterPro; IPR002341; HSF_ETS.
 DR Pfam; PF00178; Ets; 1.
 DR PRINTS; PR00454; ETSDOMAIN.
 DR SMART; SM00413; ETS; 1.
 DR PROSITE; PS00345; ETS_DOMAIN_1; 1.
 DR PROSITE; PS00345; ETS_DOMAIN_2; 1.
 DR PROSITE; PS00061; ETS_DOMAIN_3; 1.
 SQ SEQUENCE 524 AA; 56580 MW; 87834210FC7AAD1A CRC64;

Query Match 100.0%; Score 29; DB 11; Length 524;
 Best Local Similarity 100.0%; Pred. No. 4.3e+02;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 Db 98 TKPPR 92

RESULT 49

Q9VQ69 PRELIMINARY; PRT; 547 AA.
 AC Q9VQ69; Q961C7;
 DT 01-MAY-2000 (TrEMBLrel. 13, Created)
 DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE CG31672 protein (L22825p).
 GN BEST:LD15963 OR CG4248 OR CG31672.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.

```

OX NCBI_TaxID=7227;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Berkley;
RX MEDLINE=2019606; PubMed=10731132;
RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
RA Brandon R.C., Rogers Y.-H.C., Blaise R.G., Champagne M., Pfeiffer B.D.,
RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brottier P.,
RA Burris K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
RA de Pablo S.B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
RA Durbin K.J., Evangelista C.C., Ferraz C., Fertiera S., Fleischmann W.,
RA Flossler C., Gabrielian A.E., Garg N.S., Gelbart W.M., Glasser K.,
RA Glisak A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
RA Harris N.L., Harvey D., Helman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwan C.,
RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimball B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Merkulov G., Mileshina N.V., Mobarry C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Slier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wasarman D.A., Weinstock G.M., Weissbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yen R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster."
RL Science 287:2185-2195(2000).
RN [2]
RP SEQUENCE FROM N.A.
RA Celniker S.E., Adams M.D., Krommiller B., Wan K.H., Holt R.A.,
RA Evans C.A., Gocayne J.D., Amanatides P.G., Brandon R.C., Rogers Y.,
RA Banzon J., An H., Baldwin D., Banzon J., Beeson K.Y., Busam D.A.,
RA Carlson J.W., Center A., Champagne M., Davenport L.B., Dietz S.M.,
RA Dodson K., Dorsett V., Doup L.E., Doyle C., Dresnek D., Farfan D.,
RA Ferreira S., Frise E., Galle R.F., Garg N.S., George R.A.,
RA Gonzalez M., Houck J., Hoskins R.A., Hostin D., Howland T.J.,
RA Ibegwan C., Jalali M., Kruse D., Li P., Mattei B., Moshrefi A.,
RA McIntosh T.C., Moy M., Murphy B., Nelson C., Nelson K.A., Nunoo J.,
RA Pacleb J., Paragas V., Park S., Patel S., Pfeiffer B.,
RA Phouanavong S., Pittman G.S., Puri V., Richards S., Scheeler F.,
RA Stapleton M., Strong R., Svirskas R., Tector C., Tyler D.,
RA Williams S.M., Zaveri J.S., Smith H.O., Venter J.C., Rubin G.M.;
RT "Sequencing of Drosophila melanogaster genome."
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RA Misra S., Crosby M.A., Matthews B.E., Bayraktaroglu L., Campbell K.,
RA Hradecky P., Huang Y., Kanink J.S., Prochnik S.E., Smith C.D.,
RA Tupy J.L., Bergman C., Berman B., Carlson J.W., Celniker S.E.,
RA Clamp M., Drysdale R., Emmert D., Frise E., de Grey A., Harris N.,
RA Krommiller B., Marshall B., Millburn G., Richter J., Russo S.,
RA Searle S.M., Smith E., Shu S., Smutniak F., Whitfield E.,
RA Ashburner M., Gelbart W.M., Rubin G.M., Mungall C.J., Lewis S.E.;
RT "Annotation of Drosophila melanogaster genome."
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A.
RA Adams M.D., Celniker S.E., Gibbs R.A., Rubin G.M., Venter C.J.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [5]
RP SEQUENCE FROM N.A.
RA FlyBase;
RL Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
RN [6]
RP SEQUENCE FROM N.A.
RA STRAIN=Berkley;
RC Stapleton M., Brokstein P., Hong L., Agbayani A., Carlson J.,
RA Champagne M., Chavez C., Dorsett V., Farfan D., Frise E., George R.,
RA Gonzalez M., Guarin H., Li P., Liao G., Miranda A., Mungall C.J.,
RA Nunoo J., Pacleb J., Paragas V., Park S., Phouanavong S., Wan K.,
RA Yu C., Lewis S.E., Rubin G.M., Celniker S.,
RL Submitted (AUG-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AE003584; AAF51310.1; -
DR EMBL; AY051677; AAK93101.1; -
DR FlyBase; FBGN0028952; BEST:LD15963.
SQ SEQUENCE 547 AA; 63439 MW; 74CFFD0A1BA4F195 CRC64;
Query Match 100.0%; Score 29; DB 5; Length 547;
Best Local Similarity 100.0%; Pred. No. 4.4e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
Db 205 TKPPR 209
RESULT 50
Q96HX2 PRELIMINARY; PRT; 574 AA.
AC Q96HX2;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC Tissue=Eye;
RA Strausberg R.;
RL Submitted (MAY-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC007998; AAH07998.1; -
KW Hypothetical protein.
FT NON_TER 1
SQ SEQUENCE 574 AA; 60606 MW; E598819330ADFE64 CRC64;
Query Match 100.0%; Score 29; DB 4; Length 574;
Best Local Similarity 100.0%; Pred. No. 4.7e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
Db 387 TKPPR 391
Search completed: March 3, 2004, 12:18:34
Job time : 44 secs

```

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: March 3, 2004, 12:11:12 ; Search time 55 Seconds
(without alignments)
25.686 Million cell updates/sec

Title: US-09-871-974-2

Perfect score: 29

Sequence: 1 TKPRP 5 ←

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database :

1: A_Geneseq_29Jan04:*

2: Geneseq1980s:*

3: Geneseq1990s:*

4: Geneseq2000s:*

5: Geneseq2001s:*

6: Geneseq2002s:*

7: Geneseq2003as:*

8: Geneseq2003Bs:*

9: Geneseq2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	29	100.0	5	2	Aaw11052 Leukocyte
2	29	100.0	5	2	Aaw311146 Amyloid p
3	29	100.0	5	2	Aay49845 Tuftsin r
4	29	100.0	5	5	Abx08442 Tuftsin r
5	29	100.0	5	5	Abx08442 Tuftsin r
6	29	100.0	5	5	Aam51906 Tertiary
7	29	100.0	5	7	Add10684 Tuftsin a
8	29	100.0	6	2	Aaw11053 Leukocyte
9	29	100.0	6	2	Aaw311148 Amyloid p
10	29	100.0	6	2	Aaw311147 Amyloid p
11	29	100.0	6	2	Aay49840 Tuftsin r
12	29	100.0	6	5	Abx08444 Tuftsin r
13	29	100.0	6	5	Aam51904 Tertiary
14	29	100.0	6	5	Aam51905 Tertiary
15	29	100.0	6	7	Add10689 Tuftsin a
16	29	100.0	6	7	Add10687 Tuftsin a
17	29	100.0	7	2	Aay49842 Tuftsin r
18	29	100.0	7	2	Aay49841 Tuftsin r
19	29	100.0	7	5	Abx08447 Tuftsin r
20	29	100.0	7	5	Abx08445 Linear an
21	29	100.0	7	7	Add10688 Cyclic Tu
22	29	100.0	7	7	Add10686 Tuftsin a
23	29	100.0	8	2	Aar88740 Tuftsin a
24	29	100.0	8	2	Aar76218 peptide-1
25	29	100.0	8	2	Aaw11058 Leukocyte

26	29	100.0	8	2	AAV49844	Tuftsins r
27	29	100.0	8	5	ABB08448	Tuftsins r
28	29	100.0	9	2	AAR85539	Metal che
29	29	100.0	9	2	AAR85535	Metal che
30	29	100.0	9	2	AAR88735	Tuftsins a
31	29	100.0	9	2	AAR88738	Tuftsins a
32	29	100.0	9	2	AAR88741	Tuftsins a
33	29	100.0	9	2	AAR76219	Peptide-1
34	29	100.0	9	2	AAW11055	Leukocyte
35	29	100.0	9	2	AAW11059	Leukocyte
36	29	100.0	9	2	AAW11054	Leukocyte
37	29	100.0	9	2	AAW11056	Leukocyte
38	29	100.0	9	2	AAW11057	Leukocyte
39	29	100.0	9	2	AAW03420	Peptide u
40	29	100.0	9	2	AAV23752	Peptide R
41	29	100.0	10	2	AAR88739	Tuftsins a
42	29	100.0	10	7	AAR88739	Tuftsins m
43	29	100.0	10	7	AAR88736	Tuftsins m
44	29	100.0	11	2	AAR88736	Tuftsins a
45	29	100.0	11	7	ADD10691	Tuftsins a
46	29	100.0	11	7	ADD10690	Tuftsins a
47	29	100.0	12	5	ABB08446	Tuftsins r
48	29	100.0	20	5	ABB08449	99mTc rad
49	29	100.0	53	5	ABP34343	Human ORF
50	29	100.0	65	4	ABG07007	Novel hum
51	29	100.0	99	4	AAU51890	Propionib
52	29	100.0	99	5	ABB84958	Human PRO
53	29	100.0	99	5	ABG34045	Human PRO
54	29	100.0	99	5	ABB95564	Human ang
55	29	100.0	99	6	ABM48409	Propionib
56	29	100.0	99	6	ADA01300	Human PRO
57	29	100.0	99	6	ADA43729	Human sec
58	29	100.0	99	6	ADA43497	Human sec
59	29	100.0	99	6	ADA01172	Human PRO
60	29	100.0	99	7	ADA01056	Human sec
61	29	100.0	99	7	ADA43613	Human sec
62	29	100.0	99	7	ADA06875	Human PRO
63	29	100.0	99	7	ADA08363	Novel hum
64	29	100.0	99	7	ABE99656	Human PRO
65	29	100.0	99	7	ABE88939	Human PRO
66	29	100.0	99	7	ABE66094	Human sec
67	29	100.0	99	7	ABE99772	Human PRO
68	29	100.0	99	7	ABE99427	Novel hum
69	29	100.0	99	7	ABE65978	Human sec
70	29	100.0	99	7	ADC23376	Human tra
71	29	100.0	99	7	ADC28069	Human PRO
72	29	100.0	99	7	ADD10573	Human sec
73	29	100.0	99	7	ADD11533	Human sec
74	29	100.0	99	7	ADD37326	Human sec
75	29	100.0	99	7	ADE04896	Human PRO
76	29	100.0	99	7	ADE11202	Human PRO
77	29	100.0	99	7	ADD88133	Human PRO
78	29	100.0	99	7	ADD95428	Human sec
79	29	100.0	99	7	ADE06358	Human PRO
80	29	100.0	99	7	ADE38133	Human PRO
81	29	100.0	99	7	ADD88249	Human PRO
82	29	100.0	99	7	ADD90830	Human sec
83	29	100.0	99	8	ADE51683	Human sec
84	29	100.0	99	8	ADE51799	Human sec
85	29	100.0	99	8	ADE37657	Human sec
86	29	100.0	99	8	ADE37541	Human sec
87	29	100.0	99	8	ADD95312	Human sec
88	29	100.0	99	8	ADE38012	Human PRO
89	29	100.0	99	8	ADE76101	Human PRO
90	29	100.0	99	8	ADE39424	Human PRO
91	29	100.0	99	8	ADE04228	Human PRO
92	29	100.0	99	8	ADE39825	Human PRO
93	29	100.0	99	8	ADE19890	Human PRO
94	29	100.0	99	8	ADE77268	Human sec
95	29	100.0	99	8	ADE65376	Human PRO
96	29	100.0	99	8	ADE75985	Human PRO
97	29	100.0	99	8	ADE37896	Human PRO
98	29	100.0	99	8	ADE64506	Human PRO

99 29 100.0 99 8 ADE41534
100 29 100.0 99 8 ADE38841 Human PRO
101 29 100.0 99 8 ADE51915 Human sec
102 29 100.0 99 8 ADD90946 Human sec
103 29 100.0 99 8 ADE38725 Human PRO
104 29 100.0 99 8 ADE37425 Human sec
105 29 100.0 99 8 ADE06242 Human PRO
106 29 100.0 99 8 ADD90101 Human sec
107 29 100.0 99 8 ADE38609 Human PRO
108 29 100.0 99 8 ADE39540 Human PRO
109 29 100.0 99 8 ADD89145 Human PRO
110 29 100.0 99 8 ADD88912 Human PRO
111 29 100.0 99 8 ADE19806 Human PRO
112 29 100.0 99 8 ADE77384 Human sec
113 29 100.0 99 8 ADE5260 Human PRO
114 29 100.0 99 8 ADE39308 Human PRO
115 29 100.0 99 8 ADE38493 Human sec
116 29 100.0 103 4 AAU42638 Propionib
117 29 100.0 103 6 ABM39157 Propionib
118 29 100.0 107 3 AAY70535 Maize pla
119 29 100.0 136 4 ABG22236 Novel hum
120 29 100.0 149 4 ABG16691 Novel hum
121 29 100.0 198 4 AAU46263 Propionib
122 29 100.0 198 6 ABM42782 Propionib
123 29 100.0 216 4 ABG03775 Novel hum
124 29 100.0 227 4 ABG91397 Primate L
125 29 100.0 253 4 ABG07008 Novel hum
126 29 100.0 304 4 ABG16575 Novel hum
127 29 100.0 309 5 ABG77237 Selected
128 29 100.0 309 5 ABJ11108 Yeast sel
129 29 100.0 310 4 ABG06185 Novel hum
130 29 100.0 334 4 ABG17530 Novel hum
131 29 100.0 340 3 AAG43777 Arabidops
132 29 100.0 343 3 AAG43776 Arabidops
133 29 100.0 344 3 AAG43775 Arabidops
134 29 100.0 350 4 AAB50667 C. elegan
135 29 100.0 351 4 AAB50666 C. elegan
136 29 100.0 409 6 AAE34477 Murine AG
137 29 100.0 415 7 AAC774485 Sulfolobu
138 29 100.0 415 7 ADD93932 Sulfolobu
139 29 100.0 474 4 ABB59612 Drosophil
140 29 100.0 494 4 ABG03323 Novel hum
141 29 100.0 524 3 AAB03192 Rat Elf-1
142 29 100.0 530 4 ABG03809 Novel hum
143 29 100.0 530 4 ABG01390 Novel hum
144 29 100.0 563 4 ABG04846 Novel hum
145 29 100.0 563 4 ABG16158 Novel hum
146 29 100.0 576 2 AAY06477 Human tum
147 29 100.0 576 2 AAY49546 Human Nsp
148 29 100.0 576 2 AAY49545 Human Nsp
149 29 100.0 576 2 AAY49547 Human Nsp
150 29 100.0 576 2 AAY49541 Human PRO

ALIGNMENTS

RESULT 1
AAW11052
ID AAW11052 standard; peptide; 5 AA.

XX AC AAW11052;
DT 03-JUN-1997 (first entry)
XX

DE Leukocyte-targeting peptide used in diagnostic imaging.

XX Leukocyte; target; direct; chelator; radionuclide; radiolabel; isotope;
KW diagnostic imaging.

XX Synthetic.

XX WO9603427-A1.

XX 08-FEB-1996.
PD
XX 28-APR-1995; 95WC-CA000249.
PF
XX 22-JUL-1994; 94US-00279155.
PR
XX (RESO-) RESOLUTION PHARM INC.
PA
XX Pollak A, Goodbody A;
PI WPI; 1996-116994/12.
XX
XX New peptide derived radionuclide chelators and metal complexes - useful
for diagnostic imaging.
XX
XX Claim 12; Page 19; 30pp; English.
XX
XX AAW11052 is a peptide used for targeting agents to leukocytes. This
peptide can be coupled to a metal radionuclide chelator (structural
formula given in the specification), labelled with a diagnostically
useful metal isotope, to form a peptide derived radionuclide chelator.
When the chelator is coupled to a targeting molecule and labelled with a
diagnostically useful metal, it can be used to detect pathological
conditions by diagnostic imaging. For example, AAW1052 targets the
chelator to leukocytes and are useful for the rapid imaging of sites of
local inflammation. Radionuclides used include 99mTc, 64Cu, 67Cu, 97Ru,
105Rh, 109Pd, 186Re, 188Re, 198Au, 199Au, 203Pb, 212Pb and 212Bi. The
coupling of a targeting agent and radionuclide using a chelating agent is
an alternative to the direct labelling of targeting agents in which
radionuclides are typically bound at the more numerous low-affinity
sites, forming unstable complexes. The new conjugates give better
scintigraphic images in rat inflammation studies than known imaging
agents Ga-67, 99mTc-1gG, 111In-WBC and 99mTc-Nanocol. They image more
rapidly than the known agents and show superior biodistribution

XX Sequence 5 AA;
SQ

Query Match 100.0%; Score 29; DB 2; Length 5;
Best Local Similarity 100.0%; Pred. No. 1.4e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
Db 1 TKPPR 5

RESULT 2
AAW31146
ID AAW31146 standard; peptide; 5 AA.
XX AC AAW31146;
XX
XX 25-MAR-2003 (revised)
DT 23-JAN-1998 (first entry)
XX
XX Amyloid plaque-targeting peptide.
XX
XX Target; delivery; radionuclide chelator; diagnosis; therapy; detection;
KW atherosclerosis; thrombosis; Alzheimer's disease.
XX Synthetic.
XX OS
XX US5659041-A.
XX
XX 19-AUG-1997.
PD
XX 02-SEP-1994; 94US-00299636.
PF
XX 19-JUL-1993; 93US-00092911.
XX
XX (RESO-) RESOLUTION PHARM INC.
PA
XX

PI Pollak A, Kirby RA, Dunn-Dufault R;
 XX WPI; 1997-424290/39.
 XX
 XX New thioacetamyl-aminocid hydrazide compounds - useful as chemical
 PT chelator of radionuclides for radio-imaging of target tissues of
 PT diagnostic interest.
 XX
 XX
 PS Disclosure; Col 29; 20pp; English.
 XX
 XX AAW31110-W31147 are peptides used for targeting a new hydrazone-type
 CC compound to various sites of disease, e.g. atherosclerotic plaque, sites
 CC of infection, platelets, thrombus or amyloid plaque. The new compound is
 CC a radionuclide chelator and is used to radiolabel the targeting peptides
 CC for the detection and diagnostic imaging of sites of disease, e.g.
 CC amyloid plaques in Alzheimer's disease. (Updated on 25-MAR-2003 to
 CC correct PF field.)
 XX
 XX
 SQ Sequence 5 AA;
 Query Match 100.0%; Score 29; DB 2; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 Db |||||
 1 TKPPR 5
 RESULT 3
 ID AAY49845 standard; peptide; 5 AA.
 AC AAY49845;
 XX
 DT 20-JAN-2000 (first entry)
 XX
 DE Tuftsin receptor antagonist peptide.
 XX
 XX Tuftsin receptor antagonist; chelate conjugate; radiopharmaceutical;
 KW diagnosis; infection; inflammation; imaging; cancer; tumour.
 XX
 XX Synthetic.
 OS
 XX WO9951628-A1.
 PN
 XX
 PD 14-OCT-1999.
 XX
 PF 29-MAR-1999; 99WO-US006824.
 XX
 PR 03-APR-1998; 98US-0080672P.
 XX
 XX (DUPO) DU PONT PHARM CO.
 PA
 XX Edwards DS, Rajopadhye M;
 PI
 XX WPI; 1999-633729/54.
 DR
 XX
 XX New polypeptides and radiopharmaceuticals used for imaging infection,
 PT inflammation and cancer.
 PT
 XX
 PS Disclosure; Page 3; 80pp; English.
 XX
 XX The present invention describes polypeptide compounds of formula (I),
 CC capable of direct transformation into a radiopharmaceutical: Ch-Ln-(X1-X2
 CC -X3-X4-X5)d (I), where X1-X5 = amino acids; Ln = a linking group; Ch = a
 CC metal bonding unit; and d is selected from 1, 2 and 3. The
 CC radiopharmaceuticals are useful for the diagnosis of infection,
 CC inflammation and cancer. The radiopharmaceuticals bind in vivo to the
 CC tuftsin receptor on the surface of white cells which accumulate at the
 CC site of infection and inflammation and can then be detected using
 CC radiation detecting probes or by imaging using a planar or ring gamma
 CC camera. The radiopharmaceuticals can also be used in treating cancer. The

CC present sequence represents the tuftsin receptor antagonist peptide given
 CC in the present invention
 XX
 SQ Sequence 5 AA;
 Query Match 100.0%; Score 29; DB 2; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 Db |||||
 1 TKPPR 5
 RESULT 4
 ID ABB08442 standard; peptide; 5 AA.
 AC ABB08442;
 XX
 DT 01-JUL-2002 (first entry)
 XX
 DE Tuftsin receptor antagonist (TKPPR).
 XX
 XX Tuftsin; endothelial cell; drug delivery; gene therapy; NP-1;
 KW angiogenesis; tumour cell; cytostatic; antagonist.
 XX
 XX Synthetic.
 OS
 XX
 FH Key Location/Qualifiers
 FT Modified-site 1 /note= "residue optionally modified by one of the
 FT following; Thr(Obzl), DPPE-Glutaryl-
 FT Di(aminodioxoactanoyl), aminodioxoactanoyl-Thr (OBnzl),
 FT di(aminodioxoactanoyl)-Thr (OBzl), (t-Bu), Fmoc-Thr(t-Bu),
 FT H-Thr(t-Bu), Boc-Thr(But)"
 FT Modified-site 2 /note= "residue optionally modified by one of the
 FT following; Z (not further defined) or (Boc)"
 FT Modified-site 5 /note= "residue optionally modified by one of the
 FT following; OH, (NO2)Obzl, (Fmc)-t-Bu, (Fmc)-COHN"
 XX
 XX WO200191805-A2.
 PN
 XX
 PD 06-DEC-2001.
 XX
 XX 04-JUN-2001; 2001WO-US018053.
 XX
 XX 02-JUN-2000; 2000US-00585364.
 PR
 XX (BRAC) BRACCO RES USA.
 PA
 XX Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;
 PI Tweedle MF, Linder K, Nanjappan P, Raju N;
 XX
 XX WPI; 2002-195523/25.
 DR
 XX
 XX Composition for use in targeting endothelial cells, tumor cells or other
 PT cells which express NP-1 comprises a compound containing a polypeptide,
 PT linker and substrate.
 PT
 XX
 XX Claim 1; Page 2; 146pp; English.
 PS
 XX
 XX The invention relates to a composition for use in targeting endothelial
 CC cells, tumour cells, or other cells which express NP-1. The activity of
 CC compositions of the invention may be described as cytostatic. Compounds
 CC of the invention are useful in pharmaceutical compositions for inhibiting
 CC angiogenesis, for imaging and targeting an angiogenic site, endothelial
 CC cells, tumour cells or other cells that express NP-1 in a human or
 CC animal. They may also be used as ultrasound contrast agents, for staging
 CC a tumour in a human or animal, for screening for the ability of an agent
 CC to target endothelial cells, tumour cells or other cells that express NP-

CC 1. They may be used for therapeutic delivery in vivo of a bioactive agent
 CC or for treating an individual exhibiting effects of an angiogenesis or a
 CC related disorder. They may be used for delivering desirred nucleic acids
 CC to endothelial cells, tumour cells or other cells expressing NP-1, for
 CC enhancing endothelial or tumour cell-targeted gene therapy, or gene
 CC therapy targeting angiogenic cells, and for treating a human or animal
 CC with a tumour or angiogenesis-related disease. The current sequence
 CC represents the tuftsin receptor antagonist (TKPPR), of which monomers,
 CC multimers, polymers or analogues of, may be used to target endothelial
 CC cells, or cells that express markers in common with endothelial cells
 XX
 SQ Sequence 5 AA;

Query Match 100.0%; Score 29; DB 5; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06; Indels 0;
 Matches 5; Conservative 0; Mismatches 0; Gaps 0;

OY 1 TKPPR 5
 |||||
 DB 1 TKPPR 5

RESULT 5
 AAM51906
 ID AAM51906 standard; peptide; 5 AA.

XX AAM51906;

DT 01-FEB-2002 (first entry)

DE Tertiary ligand complex peptide #8.

XX Ternary ligand complex; highly functionalised phosphine ligand;
 KW disease site; cardiovascular disorder; thromboembolic disease;
 KW atherosclerosis; infectious disease; cancer; radiopharmaceutical.
 XX
 OS Synthetic.

XX Key Location/Qualifiers
 FH Modified-site 1
 FT /label= OTHER
 FT /note= "modified by 6-aminohexanamide"

XX WO200177122-A1.

PD 18-OCT-2001.

XX 06-APR-2001; 2001WO-US011387.

XX 07-APR-2000; 2000US-0195235P.

PA (DUPO) DUPONT PHARM CO.

XX Liu S;

DR WPI; 2002-010384/01.

XX New ancillary ligands are useful as ligands for radiopharmaceuticals,
 PT which are useful for e.g. radioimaging a patient, diagnosing
 PT thromboembolic disorders, atherosclerosis, infections and inflammation.

PS Example; Page 126; 210pp; English.

XX The present invention relates to novel highly functionalised phosphine
 CC ligands which can be used as ancillary ligands in radiopharmaceuticals.
 CC These radiopharmaceuticals can then be used to radioimage a patient,
 CC enabling the visualisation of platelet deposition and disease sites, and
 CC allowing the diagnosis of infection, inflammation, transplant rejection,
 CC cardiovascular diseases such as thromboembolic disorders, and cancer. The
 CC present sequence is a peptide which formed part of a ligand in the
 CC exemplification of the invention

XX Sequence 5 AA;

Query Match 100.0%; Score 29; DB 5; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06; Indels 0;
 Matches 5; Conservative 0; Mismatches 0; Gaps 0;

OY 1 TKPPR 5
 |||||
 DB 1 TKPPR 5

RESULT 6
 ADD10684
 ID ADD10684 standard; peptide; 5 AA.

XX ADD10684;

DT 01-JAN-2004 (first entry)

DE Tuftsin analogue peptide #1.

XX Phagocytosis; tuftsin; endothelial cell; inflammation; cytostatic;
 KW antiangiogenic; NP-1; ultrasound contrast agent; tumour; angiogenesis;
 KW visualisation therapy; radiotherapy.

XX Synthetic.

XX Key Location/Qualifiers
 FH Modified-site 1
 FT /label= OTHER
 FT /note= "Thr is optionally linked to an oregon green (OG)
 FT moiety"

XX US2002147136-A1.

XX 10-OCT-2002.

XX 04-JUN-2001; 2001US-00871974.

XX 02-JUN-2000; 2000US-00585364.

XX (VWRO// VON WRONSKI M A.

XX (MARI// MARINELLI E R.

XX (NUNN// NUNN A D.

XX (PILL// PILLAI R.

XX (RAMA// RAMALINGAM K.

XX (TWEE// TWEEDLE M F.

XX (LIND// LINDER K.

XX (NANG// NANJAPPAN P.

XX (RAJU// RAJU N.

XX Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;

XX Tweedle MF, Linder K, Nanjappan P, Raju N;

XX WPI; 2003-800817/75.

XX Composition used in targeting endothelial cells e.g. tumor cells
 PT comprises compounds containing monomers, multimers or polymers of L-
 PT arginine-L-threonyl-L-lysyl-L-prolyl-L-prolyl.

PS Claim 1; SEQ ID NO 2; 85pp; English.

XX The invention relates to a composition (A1) comprising compounds
 CC containing monomers, multimers or polymers of TKPPR (ADD10684).
 CC Composition (A1) comprises a compound of formula A-L-B 1, where A is the
 CC TKPPR peptide, L is a linker moiety (of formula given in the
 CC specification) and B is a substrate (or a phospholipid group,
 CC derivatisable bead attached to a fluorescent or radioactive marker,
 CC bioactive agent, delivery vehicle for genetic material, drug or
 CC therapeutic, or chelating group (preferably N 4, S 4, N 3 S, N 2 S 2 or
 CC NS 3) comprising oxa-PnAc complexed with 9m rc). The compound
 CC specifically binds to NP-1 (Vascular endothelial growth factor binding
 CC receptor transmembrane glycoprotein) or cells that express NP-1 with
 CC avidity of at least that of TKPPR. Also included are an ultrasound

CC contrast agent (c1) comprising a suspension of gas filled microbubbles
 CC comprising the TKPPR compound, an ultrasound contrast agent (c2)
 CC comprising a suspension of gas filled microballoons comprising the TKPPR
 CC compound, preparation of the TKPPR compound (which comprises conjugating
 CC the monomer, multimer or polymer of TKPPR or its analogue with a linker
 CC to obtain a compound of formula A-L, forming a covalent or non-covalent
 CC bond between A-L and the substrate B-L or forming a covalent bond between
 CC B-L and the linker to form a conjugate B-L followed by conjugation with
 CC the monomer), and a kit for preparing a radiopharmaceutical comprising
 CC the compound. The compound used for targeting endothelial cells, tumour
 CC cells or other cells which express NP-1, for inhibiting angiogenesis, for
 CC ultrasound imaging, staging a tumour, screening at least one targeted
 CC ultrasound contrast agent for the ability to target endothelial cells,
 CC tumour cells or other cells which express NP-1, for the therapeutic
 CC delivery in vivo of a bioactive agent and for delivering desired nucleic
 CC acids to endothelial cells, tumour cells or other cells which express NP-
 CC 1. The composition is also useful for visualisation therapy or
 CC radiotherapy of endothelial cells. The present sequence is the TKPPR
 CC peptide.
 CC
 CC SQ Sequence 5 AA;

Query Match 100.0%; Score 29; DB 7; Length 5;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 1 TKPPR 5

RESULT 7
 AAR85540
 ID AAR85540 standard; peptide; 6 AA.
 AC AAR85540;
 XX
 DT 17-APR-1996 (first entry)
 DE
 XX
 XX Tuftsin analogue conjugated to metal-chelating moiety.
 KW tuftsin; chelator; radiodiagnostic; imaging; leukocytes; radionuclide.
 OS Synthetic;
 XX
 XX Key Location/Qualifiers
 FH Modified-site 1
 FT /note= "the amino group of this residue is acylated with
 FT (S-Acm-mercaptopoacetyl)-Ser- (N-methyl)hydrazino-
 FT nicotinoyl"
 FT
 FN WO9503280-A1.
 XX
 PD 02-FEB-1995.
 XX
 PF 18-JUL-1994; 94WO-CA000395.
 XX
 PR 19-JUL-1993; 93US-0002911.
 XX
 XX (RESO-) RESOLUTION PHARM INC.
 PA
 XX Pollak A, Kirby RA, Dufault R;
 PI
 XX WPI; 1995-075166/10.
 DR
 XX Hydrazino-type radionuclide chelating agents - capable of conjugation to
 PT targeting molecules such as proteins, peptide(s) or antibodies and thus
 PT useful in diagnosis and therapy.
 XX
 XX Claim 29; Page 29; 35pp; English.
 PS
 CC The patent provides radionuclide chelating compounds and their
 CC conjugation products with targeting molecules such as proteins, peptides

CC or antibodies. The labelled targeting molecules may be used for in-vivo
 CC diagnosis and therapy. The chelating compound is a hydrazino-containing,
 CC N-heterocycle-containing compound, and the conjugated product is
 CC exemplified by S-Acm-mercaptopoacetyl-Ser-N-methyl-hydrazino-nicotinic
 CC acid-Gly-Thr-Lys-Pro-Arg. The peptide portion of this molecule (the
 CC present sequence) represents the tuftsin antagonist TKPPR condensed via
 CC Gly onto the chelating moiety, giving a molecule which will chelate a
 CC radionuclide such as 99mTc and target it towards leukocytes
 XX

SQ Sequence 6 AA;

Query Match 100.0%; Score 29; DB 2; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 2 TKPPR 6

RESULT 8
 AAW11053
 ID AAW11053 standard; peptide; 6 AA.
 AC AAW11053;
 XX
 DT 03-JUN-1997 (first entry)
 DE
 XX
 XX Leukocyte-targeting peptide used in diagnostic imaging.
 DE
 XX
 XX Leukocyte; target; direct; chelator; radionuclide; radiolabel; isotope;
 KW diagnostic imaging.
 XX
 XX Synthetic.
 OS
 PN WO9603427-A1.
 XX
 PD 08-FEB-1996.
 XX
 PF 28-APR-1995; 95WO-CA000249.
 XX
 PR 22-JUL-1994; 94US-00279155.
 XX
 XX (RESO-) RESOLUTION PHARM INC.
 PA
 XX Pollak A, Goodbody A;
 PI
 XX WPI; 1996-116994/12.
 DR
 XX New peptide derived radionuclide chelators and metal complexes - useful
 PT for diagnostic imaging.
 PT
 PS Claim 13; Page 20; 30pp; English.

CC AAW11053 is a peptide used for targeting agents to leukocytes. This
 CC peptide can be coupled to a metal radionuclide chelator (structural
 CC formula given in the specification) labelled with a diagnostically
 CC useful metal isotope, to form a peptide derived radionuclide chelator.
 CC When the chelator is coupled to a targeting molecule and labelled with a
 CC diagnostically useful metal, it can be used to detect pathological
 CC conditions by diagnostic imaging. For example, AAW11053 targets the
 CC chelator to leukocytes and are useful for the rapid imaging of sites of
 CC local inflammation. Radionuclides used include 99mTc, 64Cu, 67Cu, 97Ru,
 CC 105Rh, 109Pd, 186Re, 188Re, 198Au, 203Pb, 212Pb and 212Bi. The
 CC coupling of a targeting agent and radionuclide using a chelating agent is
 CC an alternative to the direct labelling of targeting agents in which
 CC radionuclides are typically bound at the more numerous low-affinity
 CC sites, forming unstable complexes. The new conjugates give better
 CC scintigraphic images in rat inflammation studies than known imaging
 CC agents Ga-67, 99mTc-IGG, 111In-WBC and 99mTc-Nanocol. They image more
 CC rapidly than the known agents and show superior biodistribution
 XX
 XX Sequence 6 AA;

Query Match 100.0%; Score 29; DB 2; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TKPPR 5
 Db 2 TKPPR 6

RESULT 9
 AAW31148
 ID AAW31148 standard; peptide; 6 AA.
 XX
 AC AAW31148;
 XX
 XX 25-MAR-2003 (revised)
 DT 23-JAN-1998 (first entry)
 XX
 DE Amyloid plaque-targeting peptide-radiolabeled chelator conjugate.
 XX
 XX Target; delivery; radiolabeled chelator; diagnosis; therapy; detection;
 KW atherosclerosis; thrombosis; Alzheimer's disease.
 XX
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 FT Modified-site 1
 FT /note= "N terminally acetylated with S-Acm-Mercaptoacetyl
 FT -Ser-N- methylhydrazino nicotinic acid"
 XX
 XX US5659041-A.
 PN
 XX 19-AUG-1997.
 PD
 XX 02-SEP-1994; 94US-00299636.
 PF
 XX 19-JUL-1993; 93US-00092911.
 PR
 XX (RESO-) RESOLUTION PHARM INC.
 PA
 XX Pollak A, Kirby RA, Dunn-Dufault R;
 PI WPI; 1997-424290/39.
 DR
 XX New thio:acetyl-aminoacid hydrazide compounds - useful as chemical
 PT chelator of radionuclides for radio:imaging of target tissues of
 PT diagnostic interest.
 XX
 XX Example 11; Col 14; 20pp; English.
 PS
 XX AAW31148 is a peptide conjugate of an amyloid plaque-targeting peptide
 CC and a new hydrazino-type compound. The new compound is a radionuclide
 CC chelator and is used to radiolabel targeting peptides for the detection
 CC and diagnostic imaging of sites of disease, e.g. amyloid plaques in
 CC Alzheimer's disease, thrombi in thrombosis or other sites of infection.
 CC (Updated on 25-MAR-2003 to correct PF field.)
 XX
 SQ Sequence 6 AA;
 Query Match 100.0%; Score 29; DB 2; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TKPPR 5
 Db 2 TKPPR 6

RESULT 10
 AAW31147
 ID AAW31147 standard; peptide; 6 AA.
 XX
 AC AAW31147;
 XX
 XX 25-MAR-2003 (revised)
 DT 23-JAN-1998 (first entry)
 XX
 DE Amyloid plaque-targeting peptide.
 XX
 XX Target; delivery; radionuclide chelator; diagnosis; therapy; detection;
 KW atherosclerosis; thrombosis; Alzheimer's disease.
 XX
 OS Synthetic.
 XX
 XX US5659041-A.
 PN
 XX 19-AUG-1997.
 PD
 XX 02-SEP-1994; 94US-00299636.
 PF
 XX 19-JUL-1993; 93US-00092911.
 PR
 XX (RESO-) RESOLUTION PHARM INC.
 PA
 XX Pollak A, Kirby RA, Dunn-Dufault R;
 PI WPI; 1997-424290/39.
 DR
 XX New thio:acetyl-aminoacid hydrazide compounds - useful as chemical
 PT chelator of radionuclides for radio:imaging of target tissues of
 PT diagnostic interest.
 XX
 XX Disclosure; Col 29; 20pp; English.
 PS
 XX AAW3110-W31147 are peptides used for targeting a new hydrazino-type
 CC compound to various sites of disease, e.g. atherosclerotic plaque, sites
 CC of infection, platelets, thrombus or amyloid plaque. The new compound is
 CC a radionuclide chelator and is used to radiolabel the targeting peptides
 CC for the detection and diagnostic imaging of sites of disease, e.g.
 CC amyloid plaques in Alzheimer's disease. (Updated on 25-MAR-2003 to
 CC correct PF field.)
 XX
 SQ Sequence 6 AA;
 Query Match 100.0%; Score 29; DB 2; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TKPPR 5
 Db 2 TKPPR 6

RESULT 11
 AAY49840
 ID AAY49840 standard; peptide; 6 AA.
 XX
 AC AAY49840;
 XX
 XX 20-JAN-2000 (first entry)
 DT
 XX
 DE Tuftsin receptor antagonist chelate conjugate #1.
 XX
 XX Tuftsin receptor antagonist; chelate conjugate; radiopharmaceutical;
 KW diagnosis; infection; inflammation; imaging; cancer; tumour.
 XX
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 FT Modified-site 1
 FT /label= Acp
 FT /note= "modified 6-aminocaproic acid: 6-((6-((1-aza-2-
 FT -sulphophenyl)vinyl)amino)-3-pyridyl)
 FT carbonylamino)hexanoyl"
 FT
 XX

PN WO9951628-A1.
 XX
 PD 14-OCT-1999.
 XX
 PF 29-MAR-1999; 99WO-US006824.
 XX
 PR 03-APR-1998; 98US-0080672P.
 XX
 PA (DUPO) DU PONT PHARM CO.
 XX
 XX Edwards DS, Rajopadhye M;
 PI WPI; 1999-633729/54.
 XX
 DR New polypeptides and radiopharmaceuticals used for imaging infection,
 XX inflammation and cancer.
 PT
 PT
 XX
 PS Claim 4; Page 71; 80pp; English.
 XX
 CC The present invention describes polypeptide compounds of formula (I),
 CC capable of direct transformation into a radiopharmaceutical: $\text{Ch-Ln-(X1-X2-X3-X4-X5)d}$ (I), where X1-X5 = amino acids; Ln = a linking group; Ch = a
 CC metal bonding unit; and d is selected from 1, 2 and 3. The
 CC radiopharmaceuticals are useful for the diagnosis of infection,
 CC inflammation and cancer. The radiopharmaceuticals bind in vivo to the
 CC tuftsin receptor on the surface of white cells which accumulate at the
 CC site of infection and inflammation and can then be detected using
 CC radiation detecting probes or by imaging using a planar or ring gamma
 CC camera. The radiopharmaceuticals can also be used in treating cancer. The
 CC present sequence represents a specifically claimed tuftsin receptor
 CC antagonist chelate conjugate from the present invention
 CC
 XX
 SQ Sequence 6 AA;
 Query Match 100.0%; Score 29; DB 2; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 2 TKPPR 6
 RESULT 12
 ABB08444
 ID ABB08444 standard; peptide; 6 AA.
 AC ABB08444;
 XX
 XX 01-JUL-2002 (first entry)
 DT
 DE Tuftsin receptor antagonist (TKPPR) derivative peptide 2.
 XX
 XX Tuftsin; endothelial cell; drug delivery; gene therapy; NP-1;
 KW angiogenesis; tumour cell; cytostatic; antagonist.
 KW
 XX Synthetic.
 OS
 XX
 XX Key Location/Qualifiers
 FH Modified-site 1
 FT /note= "residue optionally modified by one of the
 FT following; DPPG-Glutaryl, N-Glutaryl-Gly, NH2, Oregon
 FT Green (5 isomer), Fmoc"
 FT Modified-site 2
 FT /note= "residue optionally modified by one of the
 FT following; Obz1, (tBu)"
 FT Modified-site 3
 FT /note= "residue optionally modified by one of the
 FT following; 2 (not further defined), (Mtc), (Boc)"
 FT Modified-site 6
 FT /note= "residue optionally modified by one of the
 FT following; OH, (NO2)Obz1, (NO2), (Pmc)Obz1, (pmc)-wang-
 FT resin"

XX WO200191805-A2.
 XX
 XX 06-DEC-2001.
 XX
 XX 04-JUN-2001; 2001WO-US018053.
 XX
 XX 02-JUN-2000; 2000US-00585364.
 XX
 XX (BRAC) BRACCO RES USA.
 XX
 XX Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;
 PI Tweedle MF, Linder K, Nanjappan P, Raju N;
 XX WPI; 2002-195523/25.
 DR
 XX
 XX Composition for use in targeting endothelial cells, tumor cells or other
 PT cells which express NP-1 comprises a compound containing a polypeptide,
 PT linker and substrate.
 XX
 XX Example 4; Page 67; 145pp; English.
 XX
 CC The invention relates to a composition for use in targeting endothelial
 CC cells, tumour cells, or other cells which express NP-1. The activity of
 CC compositions of the invention may be described as cytostatic. Compounds
 CC of the invention are useful in pharmaceutical compositions for inhibiting
 CC angiogenesis, for imaging and targeting an angiogenic site, endothelial
 CC cells, tumour cells or other cells that express NP-1 in a human or
 CC animal. They may also be used as ultrasound contrast agents, for staging
 CC a tumour in a human or animal, for screening for the ability of an agent
 CC to target endothelial cells, tumour cells or other cells that express NP-1.
 CC 1. They may be used for therapeutic delivery in vivo of a bioactive agent
 CC or for treating an individual exhibiting effects of an angiogenesis or a
 CC related disorder. They may be used for delivering desired nucleic acids
 CC to endothelial cells, tumour cells or other cells expressing NP-1, for
 CC enhancing endothelial or tumour cell-targeted gene therapy, or gene
 CC therapy targeting angiogenic cells, and for treating a human or animal
 CC with a tumour or angiogenesis-related disease. The current sequence
 CC represents a tuftsin receptor antagonist (TKPPR) derivative of the
 CC invention
 XX
 SQ Sequence 6 AA;
 Query Match 100.0%; Score 29; DB 5; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 2 TKPPR 6
 RESULT 13
 AAM51904
 ID AAM51904 standard; peptide; 6 AA.
 XX
 XX AAM51904;
 AC
 XX
 XX 01-FEB-2002 (first entry)
 DT
 DE Tertiary ligand complex peptide #6.
 XX
 XX Ternary ligand complex; highly functionalised phosphine ligand;
 KW disease site; cardiovascular disorder; thromboembolic disease;
 KW atherosclerosis; infectious disease; cancer; radiopharmaceutical.
 KW
 XX Synthetic.
 OS
 XX
 XX Key Location/Qualifiers
 FH Misc-difference 1
 FT /note= "D-form residue"
 FT Modified-site 2
 FT /label= OTHER

/note= "modified by 6-aminohexanamide"

PA (DUPO) DUPONT PHARM CO.
 XX
 PI Liu S;
 XX
 DR WPI; 2002-010884/01.
 XX
 PT New ancillary ligands are useful as ligands for radiopharmaceuticals,
 which are useful for e.g. radioimaging a patient, diagnosing
 PT thromboembolic disorders, atherosclerosis, infections and inflammation.
 XX
 PS Example; Page 125; 210pp; English.
 XX
 CC The present invention relates to novel highly functionalised phosphine
 CC ligands which can be used as ancillary ligands in radiopharmaceuticals.
 CC These radiopharmaceuticals can then be used to radioimage a patient,
 CC enabling the visualisation of platelet deposition and disease sites, and
 CC allowing the diagnosis of infection, inflammation, transplant rejection,
 CC cardiovascular diseases such as thromboembolic disorders, and cancer. The
 CC present sequence is a peptide which formed part of a ligand in the
 CC exemplification of the invention
 XX
 SQ Sequence 6 AA;

Query Match 100.0%; Score 29; DB 5; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 2 TKPPR 6

RESULT 15

ADD10689
 ID ADD10689 standard; peptide; 6 AA.

XX
 AC ADD10689;
 XX
 DT 01-JAN-2004 (first entry)
 XX
 DE Tuftsin analogue peptide #4.

XX Phagocytosis; tuftsin; endothelial cell; inflammation; cytostatic;
 KW antiangiogenic; NP-1; ultrasound contrast agent; tumour; angiogenesis;
 KW visualisation therapy; radiotherapy.

XX Synthetic.

XX Key Location/Qualifiers
 FT Modified-site 1 /label= OTHER
 FT /note= "Glu is covalently linked to a DPPE
 (dipalmitoylphosphatidylethanolamine) moiety"
 FT Modified-site 2
 FT /label= OTHER
 FT /note= "Thr is di(aminodioxanoyl)-Thr"

US2002147136-A1.

PD 10-OCT-2002.

XX 04-JUN-2001; 2001US-00871974.

XX 02-JUN-2000; 2000US-00585364.

XX (VWRO/) VON WRONSKI M A.

XX (MARI/) MARINELLI E R.

XX (NUNN/) NUNN A D.

XX (PILL/) PILLAI R.

XX (RAMA/) RAMALINGAM K.

XX (TWE/) TWEEDLE M F.

XX (LIND/) LINDER K.

XX (NANJ/) NANJAPPAN P.

FT
 XX WO200177122-A1.
 PN
 XX 18-OCT-2001.
 PD

XX 06-APR-2001; 2001WO-US011387.

XX 07-APR-2000; 2000US-0195235P.

XX (DUPO) DUPONT PHARM CO.

XX Liu S;

XX WPI; 2002-010884/01.

XX New ancillary ligands are useful as ligands for radiopharmaceuticals,
 PT which are useful for e.g. radioimaging a patient, diagnosing
 PT thromboembolic disorders, atherosclerosis, infections and inflammation.
 XX
 PS Example; Page 125; 210pp; English.

XX The present invention relates to novel highly functionalised phosphine
 CC ligands which can be used as ancillary ligands in radiopharmaceuticals.
 CC These radiopharmaceuticals can then be used to radioimage a patient,
 CC enabling the visualisation of platelet deposition and disease sites, and
 CC allowing the diagnosis of infection, inflammation, transplant rejection,
 CC cardiovascular diseases such as thromboembolic disorders, and cancer. The
 CC present sequence is a peptide which formed part of a ligand in the
 CC exemplification of the invention
 XX
 SQ Sequence 6 AA;

Query Match 100.0%; Score 29; DB 5; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 2 TKPPR 6

RESULT 14

AAM51905
 ID AAM51905 standard; peptide; 6 AA.

XX
 AC AAM51905;

XX 01-FEB-2002 (first entry)

XX Tertiary ligand complex peptide #7.

XX Ternary ligand complex; highly functionalised phosphine ligand;
 KW disease site; cardiovascular disorder; thromboembolic disease;
 KW atherosclerosis; infectious disease; cancer; radiopharmaceutical.

XX Synthetic.

XX Key Location/Qualifiers

FT Misc-difference 1 /note= "D-form residue"

FT Modified-site 2

FT /label= OTHER

FT /note= "modified by 6-aminohexanamide"

XX WO200177122-A1.

XX 18-OCT-2001.

XX 06-APR-2001; 2001WO-US011387.

XX 07-APR-2000; 2000US-0195235P.

PA (RAJU/) RAJU N.
XX Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;
PI Tweedle MF, Linder K, Nanjappan P, Raju N;
XX WPI; 2003-800817/75.
XX
XX Composition used in targeting endothelial cells e.g. tumor cells
PT comprises compounds containing monomers, multimers or polymers of L-
PT arginine-L-threonyl-L-lysyl-L-prolyl-L-prolyl.
XX
XX Example 23; Page 44; 85pp; English.
XX
XX The invention relates to a composition (A1) comprising compounds
CC containing monomers, multimers or polymers of TKPPR (ADD10684).
CC Composition (A1) comprises a compound of formula A-L-B 1, where A is the
CC TKPPR peptide, L is a linker moiety (of formula given in the
CC specification), and B is a substrate (or a phospholipid group,
CC derivatisable bead attached to a fluorescent or radioactive marker,
CC bioactive agent, delivery vehicle for genetic material, drug or
CC therapeutic, or chelating group (preferably N 4, S 4, N 3 S, N 2 S 2 or
CC NS 3) comprising oxa-PnAO complexed with 99m Tc). The compound
CC specifically binds to NP-1 (Vascular endothelial growth factor binding
CC receptor transmembrane glycoprotein) or cells that express NP-1 with
CC avidity of at least that of TKPPR. Also included are an ultrasound
CC contrast agent (c1) comprising a suspension of gas filled microbubbles
CC comprising the TKPPR compound, an ultrasound contrast agent (c2)
CC comprising a suspension of gas filled microballoons comprising the TKPPR
CC compound, preparation of the TKPPR compound (which comprises conjugating
CC the monomer, multimer or polymer of TKPPR or its analogue with a linker
CC to obtain a compound of formula A-L, forming a covalent or non-covalent
CC bond between A-L and the substrate B 1 or forming a covalent bond between
CC B 1 and the linker to form a conjugate B-L followed by conjugation with
CC the monomer), and a kit for preparing a radiopharmaceutical comprising
CC the compound. The compound used for targeting endothelial cells, tumour
CC cells or other cells which express NP-1, for inhibiting angiogenesis, for
CC ultrasound imaging, staging a tumour, screening at least one targeted
CC ultrasound contrast agent for the ability to target endothelial cells,
CC tumour cells or other cells which express NP-1, for the therapeutic
CC delivery in vivo of a bioactive agent and for delivering desired nucleic
CC acids to endothelial cells, tumour cells or other cells which express NP-
CC 1. The composition is also useful for visualisation therapy or
CC radiotherapy of endothelial cells. The present sequence is a TKPPR
CC analogue peptide.
XX
XX Sequence 6 AA;
XX
XX Query Match 100.0%; Score 29; DB 7; Length 6;
XX Best Local Similarity 100.0%; Pred. No. 1.4e+06;
XX Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 TKPPR 5
XX |||||
XX 2 TKPPR 6
XX
XX RESULT 16
XX ADD10687
XX ID ADD10687 standard; peptide; 6 AA.
XX AC ADD10687;
XX XX
XX 01-JAN-2004 (first entry)
XX
XX Tuftsin analogue peptide #3.
XX
XX Phagocytosis; tuftsin; endothelial cell; inflammation; cytostatic;
XX antiangiogenic; NP-1; ultrasound contrast agent; tumour; angiogenesis;
XX visualisation therapy; radiotherapy.
XX Synthetic.
XX
XX Key Location/Qualifiers
XX FH

FT Modified-site 1 /label= OTHER
FT FT
FT FT
XX XX
XX PN US2002147136-A1.
XX
XX PD 10-OCT-2002.
XX
XX PF 04-JUN-2001; 2001US-00871974.
XX
XX PR 02-JUN-2000; 2000US-00585364.
XX
XX (VWRO/) VON WRONSKI M A.
XX PA (MARI/) MARINELLI E R.
XX PA (NUNN/) NUNN A D.
XX PA (PILL/) PILLAI R.
XX PA (RAMA/) RAMALINGAM K.
XX PA (TWEE/) TWEDDLE M F.
XX PA (LIND/) LINDER K.
XX PA (NANJ/) NANJAPPAN P.
XX PA (RAJU/) RAJU N.
XX
XX Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;
PI Tweedle MF, Linder K, Nanjappan P, Raju N;
XX WPI; 2003-800817/75.
XX
XX Composition used in targeting endothelial cells e.g. tumor cells
PT comprises compounds containing monomers, multimers or polymers of L-
PT arginine-L-threonyl-L-lysyl-L-prolyl-L-prolyl.
XX
XX Example 14; Page 38; 85pp; English.
XX
XX The invention relates to a composition (A1) comprising compounds
CC containing monomers, multimers or polymers of TKPPR (ADD10684).
CC Composition (A1) comprises a compound of formula A-L-B 1, where A is the
CC TKPPR peptide, L is a linker moiety (of formula given in the
CC specification), and B is a substrate (or a phospholipid group,
CC derivatisable bead attached to a fluorescent or radioactive marker,
CC bioactive agent, delivery vehicle for genetic material, drug or
CC therapeutic, or chelating group (preferably N 4, S 4, N 3 S, N 2 S 2 or
CC NS 3) comprising oxa-PnAO complexed with 99m Tc). The compound
CC specifically binds to NP-1 (Vascular endothelial growth factor binding
CC receptor transmembrane glycoprotein) or cells that express NP-1 with
CC avidity of at least that of TKPPR. Also included are an ultrasound
CC contrast agent (c1) comprising a suspension of gas filled microbubbles
CC comprising the TKPPR compound, an ultrasound contrast agent (c2)
CC comprising a suspension of gas filled microballoons comprising the TKPPR
CC compound, preparation of the TKPPR compound (which comprises conjugating
CC the monomer, multimer or polymer of TKPPR or its analogue with a linker
CC to obtain a compound of formula A-L, forming a covalent or non-covalent
CC bond between A-L and the substrate B 1 or forming a covalent bond between
CC B 1 and the linker to form a conjugate B-L followed by conjugation with
CC the monomer), and a kit for preparing a radiopharmaceutical comprising
CC the compound. The compound used for targeting endothelial cells, tumour
CC cells or other cells which express NP-1, for inhibiting angiogenesis, for
CC ultrasound imaging, staging a tumour, screening at least one targeted
CC ultrasound contrast agent for the ability to target endothelial cells,
CC tumour cells or other cells which express NP-1, for the therapeutic
CC delivery in vivo of a bioactive agent and for delivering desired nucleic
CC acids to endothelial cells, tumour cells or other cells which express NP-
CC 1. The composition is also useful for visualisation therapy or
CC radiotherapy of endothelial cells. The present sequence is a TKPPR
CC analogue peptide.
XX
XX Sequence 6 AA;
XX
XX Query Match 100.0%; Score 29; DB 7; Length 6;
XX Best Local Similarity 100.0%; Pred. No. 1.4e+06;
XX Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 TKPPR 5
XX |||||
XX 2 TKPPR 6
XX
XX RESULT 16
XX ADD10687
XX ID ADD10687 standard; peptide; 6 AA.
XX AC ADD10687;
XX XX
XX 01-JAN-2004 (first entry)
XX
XX Tuftsin analogue peptide #3.
XX
XX Phagocytosis; tuftsin; endothelial cell; inflammation; cytostatic;
XX antiangiogenic; NP-1; ultrasound contrast agent; tumour; angiogenesis;
XX visualisation therapy; radiotherapy.
XX Synthetic.
XX
XX Key Location/Qualifiers
XX FH

```
Db          |||||
            2 TKPPR 6

RESULT 17
AAV49842
ID AAV49842 standard; peptide; 7 AA.
XX
XX AAV49842;
AC
XX
XX 20-JAN-2000 (first entry)
XX
XX Tuftsin receptor antagonist chelate conjugate #3.
DE
XX
XX Tuftsin receptor antagonist; chelate conjugate; radiopharmaceutical;
KW diagnosis; infection; inflammation; imaging; cancer; tumour.
KW
XX
XX Synthetic.
OS
XX
XX Key Location/Qualifiers
FH Modified-site 1
FT /note= "modified tyrosine: ((6-((1-aza-2- (2-
FT sulphophenyl)vinyl)amino)-3-pyridyl) carbonyl)-L-
FT phenylalanine"
FT Modified-site 2
FT /label= Acp
FT
FT WO9951628-A1.
XX
XX PN
XX 14-OCT-1999.
XX
XX PD
XX 29-MAR-1999; 99WO-US006824.
XX
XX PF
XX 03-APR-1998; 98US-0080672P.
XX
XX PR
XX (DUPO ) DU PONT PHARM CO.
XX
XX PA
XX Edwards DS, Rajopadhye M;
XX
XX PI
XX WPI; 1999-633729/54.
XX
XX DR
XX New polypeptides and radiopharmaceuticals used for imaging infection,
XX inflammation and cancer.
XX
XX PS Claim 4; Page 71; 80pp; English.
XX
XX CC The present invention describes polypeptide compounds of formula (I),
XX capable of direct transformation into a radiopharmaceutical: Ch-Ln-(X1-X2
XX -X3-X4-X5)d (I), where X1-X5 = amino acids; Ln = a linking group; Ch = a
XX metal bonding unit; and d is selected from 1, 2 and 3. The
XX radiopharmaceuticals are useful for the diagnosis of infection,
XX inflammation and cancer. The radiopharmaceuticals bind in vivo to the
XX tuftsin receptor on the surface of white cells which accumulate at the
XX site of infection and inflammation and can then be detected using
XX radiation detecting probes or by imaging using a planar or ring gamma
XX camera. The radiopharmaceuticals can also be used in treating cancer. The
XX present sequence represents a specifically claimed tuftsin receptor
XX antagonist chelate conjugate from the present invention
XX
XX SQ Sequence 7 AA;
XX Query Match 100.0%; Score 29; DB 2; Length 7;
XX Best Local Similarity 100.0%; Pred. No. 1.4e+06;
XX Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 TKPPR 5
XX |||||
XX 3 TKPPR 7
XX
XX Db
XX
XX RESULT 18
XX AAV49841
XX ID AAV49841 standard; peptide; 7 AA.
XX
```


Db 2 TKPPR 6

RESULT 21
ADD10688
ID ADD10688 standard; peptide; 7 AA.
AC ADD10688;
XX
DT 01-JAN-2004 (first entry)
XX
DE Cyclic Tuftsin analogue peptide.
XX
KW Phagocytosis; tuftsin; endothelial cell; inflammation; cytostatic;
KW antiangiogenic; NP-1; ultrasound contrast agent; tumour; angiogenesis;
KW visualisation therapy; radiotherapy; cyclic.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Disulfide-bond 1..7
FT /note= "The molecule is cyclised via this disulfide bond"
PN
PN US2002147136-A1.
XX
XX 10-OCT-2002.
XX
XX 04-JUN-2001; 2001US-00871974.
XX
XX 02-JUN-2000; 2000US-00585364.
XX
XX (VWRO/) VON WRONSKI M A.
XX (WARI/) MARINELLI E R.
XX (NUNN/) NUNN A D.
XX (PILL/) PILLAI R.
XX (RAMA/) RAMALINGAM K.
XX (TWEE/) TWEEDELE M F.
XX (LIND/) LINDER K.
XX (NANJ/) NANJAPPAN P.
XX (RAJU/) RAJU N.
XX
XX Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;
XX Tweedle MF, Linder K, Nanjappan P, Raju N;
XX WPI; 2003-800817/75.
XX
XX Composition used in targeting endothelial cells e.g. tumor cells
XX comprises compounds containing monomers, multimers or polymers of L-
XX arginine-L-threonine-L-lysyl-L-prolyl-L-prolyl.
XX
XX Example 21; Page 41; 85pp; English.
XX
XX The invention relates to a composition (A1) comprising compounds
XX containing monomers, multimers or polymers of TKPPR (ADD10684).
XX Composition (A1) comprises a compound of formula A-L-B 1, where A is the
XX TKPPR peptide, L is a linker moiety (of formula given in the
XX specification) and B is a substrate (or a phospholipid group,
XX derivatisable bead attached to a fluorescent or radioactive marker,
XX bioactive agent, delivery vehicle for genetic material, drug or
XX therapeutic, or chelating group (preferably N 4, S 4, N 3 S, N 2 S 2 or
XX NS 3) comprising oxa-TnAO complexed with 99m Tc). The compound
XX specifically binds to NP-1 (Vascular endothelial growth factor binding
XX receptor transmembrane glycoprotein) or cells that express NP-1 with
XX avidity of at least that of TKPPR. Also included are an ultrasound
XX contrast agent (c1) comprising a suspension of gas filled microbubbles
XX comprising the TKPPR compound, an ultrasound contrast agent (c2)
XX comprising a suspension of gas filled microballoons comprising the TKPPR
XX compound, preparation of the TKPPR compound (which comprises conjugating
XX the monomer, multimer or polymer of TKPPR or its analogue with a linker
XX to obtain a compound of formula A-L, forming a covalent or non-covalent
XX bond between A-L and the substrate B 1 or forming a covalent bond between
XX B 1 and the linker to form a conjugate B-L followed by conjugation with

CC the monomer), and a kit for preparing a radiopharmaceutical comprising
CC the compound. The compound used for targeting endothelial cells, tumour
CC cells or other cells which express NP-1, for inhibiting angiogenesis, for
CC ultrasound imaging, staging a tumour, screening at least one targeted
CC ultrasound contrast agent for the ability to target endothelial cells,
CC tumour cells or other cells which express NP-1, for the therapeutic
CC delivery in vivo of a bioactive agent and for delivering desired nucleic
CC acids to endothelial cells, tumour cells or other cells which express NP-
CC 1. The composition is also useful for visualisation therapy or
CC radiotherapy of endothelial cells. The present sequence is a cyclic TKPPR
XX analogue peptide.
XX
XX Sequence 7 AA;
XX
XX Query Match 100.0%; Score 29; DB 7; Length 7;
XX Best Local Similarity 100.0%; Pred. No. 1.4e+06;
XX Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 1 TKPPR 5
Db 2 TKPPR 6

RESULT 22
ADD10686
ID ADD10686 standard; peptide; 7 AA.
XX
XX ADD10686;
XX
DT 01-JAN-2004 (first entry)
XX
XX Tuftsin analogue peptide #2.
XX
XX Phagocytosis; tuftsin; endothelial cell; inflammation; cytostatic;
XX antiangiogenic; NP-1; ultrasound contrast agent; tumour; angiogenesis;
XX visualisation therapy; radiotherapy.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX Modified-site 1
XX /label= OTHER
XX /note= "Glu is covalently linked to a DPPE
XX (dipalmitoylphosphatidylethanolamine) moiety"
XX
XX US2002147136-A1.
XX
XX 10-OCT-2002.
XX
XX 04-JUN-2001; 2001US-00871974.
XX
XX 02-JUN-2000; 2000US-00585364.
XX
XX (VWRO/) VON WRONSKI M A.
XX (WARI/) MARINELLI E R.
XX (NUNN/) NUNN A D.
XX (PILL/) PILLAI R.
XX (RAMA/) RAMALINGAM K.
XX (TWEE/) TWEEDELE M F.
XX (LIND/) LINDER K.
XX (NANJ/) NANJAPPAN P.
XX (RAJU/) RAJU N.
XX
XX Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;
XX Tweedle MF, Linder K, Nanjappan P, Raju N;
XX WPI; 2003-800817/75.
XX
XX Composition used in targeting endothelial cells e.g. tumor cells
XX comprises compounds containing monomers, multimers or polymers of L-
XX arginine-L-threonine-L-lysyl-L-prolyl-L-prolyl.
XX
XX Example 21; Page 34; 85pp; English.

XX The invention relates to a composition (A1) comprising compounds
 CC containing monomers, multimers or polymers of TKPPR (ABD10684).
 CC composition (A1) comprises a compound of formula A-L-B 1, where A is the
 CC TKPPR peptide, L is a linker moiety (cf formula given in the
 CC specification) and B is a substrate (or a phospholipid group,
 CC derivatisable bead attached to a fluorescent or radioactive marker,
 CC bioactive agent, delivery vehicle for genetic material, drug or
 CC therapeutic, or chelating group (preferably N 4, S 4, N 3 S, N 2 S 2 or
 CC NS 3) comprising oxo-PnAO complexed with 99mTc). The compound
 CC specifically binds to NP-1 (Vascular endothelial growth factor binding
 CC receptor transmembrane glycoprotein) or cells that express NP-1 with
 CC avidity of at least that of TKPPR. Also included are an ultrasound
 CC contrast agent (C1) comprising a suspension of gas filled microbubbles
 CC comprising the TKPPR compound, an ultrasound contrast agent (C2)
 CC comprising a suspension of gas filled microbubbles comprising the TKPPR
 CC compound, preparation of the TKPPR compound (which comprises conjugating
 CC the monomer, multimer or polymer of TKPPR or its analogue with a linker
 CC to obtain a compound of formula A-L, forming a covalent or non-covalent
 CC bond between A-L and the substrate B 1 or forming a covalent bond between
 CC B 1 and the linker to form a conjugate B-L followed by conjugation with
 CC the monomer), and a kit for preparing a radiopharmaceutical comprising
 CC the compound, and the compound used for targeting endothelial cells, tumour
 CC cells or other cells which express NP-1, for inhibiting angiogenesis, for
 CC ultrasound imaging, staging a tumour, screening at least one targeted
 CC ultrasound contrast agent for the ability to target endothelial cells,
 CC tumour cells or other cells which express NP-1, for the therapeutic
 CC delivery in vivo of a bioactive agent and for delivering desired nucleic
 CC acids to endothelial cells, tumour cells or other cells which express NP-
 CC 1. The composition is also useful for visualisation therapy or
 CC radiotherapy of endothelial cells. The present sequence is a TKPPR
 CC analogue peptide.

XX Sequence 7 AA;

Query Match 100.0%; Score 29; DB 7; Length 7;
 Best Local Similarity 100.0%; Pred. NO. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 DB 3 TKPPR 7

RESULT 23

AAR88740
 ID AAR88740 standard; peptide; 8 AA.

XX AAR88740;

AC AAR88740;

DT 10-APR-1996 (first entry)

DE Tuftsin antagonist peptide-metal chelator conjugate.

XX Peptide-chelator conjugate; metal chelator; diagnostic imaging;

KW inflammation; radionuclide; tuftsin; analogue; antagonist.

OS Synthetic.

XX Key Location/Qualifiers

FT Peptide 1..5

FT Modified-site 6 /label= tuftsin_antagonist

FT /label= bala

FT /note= "linking group"

FT Modified-site 7

FT /label= OTHER

FT /note= "the side-chain (epsilon) amino group of Lys at

FT position 7 forms a peptide bond with the C-terminus of

FT the tetrapeptide N',N'-dimethylglycyl-Ser-Cys(Acm)-Gly-;

FT this sidechain group consists of a tripeptide metal

FT chelator and a linking residue (Gly)"

PN WO9522996-A2.

XX 31-AUG-1995.

XX 24-FEB-1995; 95WO-CA000106.

XX 25-FEB-1994; 94US-00202178.

XX (RESO-) RESOLUTION PHARM INC.

XX Goodbody A, Pollak A;

XX WPI; 1995-311386/40.

XX New peptide-chelator conjugate and complex with traceable metal - used to
 PT image sites of inflammation in vivo without significant accumulation on
 PT the gastrointestinal tract.

XX Claim 19; Page 21; 23pp; English.

XX The present sequence is that of a specifically claimed peptide- chelator
 CC conjugate in which a tuftsin antagonist peptide is coupled to a metal
 CC chelator, via a linking group. The chelator serves as a labelling site
 CC for radionuclide metals such as technetium-99m. The tuftsin antagonist
 CC targets the conjugate to macrophages and neutrophils at sites of
 CC inflammation without significant accumulation in the gastrointestinal
 CC tract (unlike the native tuftsin tetrapeptide). The conjugate is thus
 CC useful for diagnostic imaging of inflammation sites, providing an
 CC improved target to background ratio

XX Sequence 8 AA;

Query Match 100.0%; Score 29; DB 2; Length 8;

Best Local Similarity 100.0%; Pred. NO. 1.4e+06;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5

DB 1 TKPPR 5

RESULT 24

AAR76218
 ID AAR76218 standard; peptide; 8 AA.

XX AAR76218;

XX 12-JAN-1996 (first entry)

XX peptide-ligand conjugate #1.

XX Peptide-ligand; metal-labelled imaging agent; maleimide; metal atom;
 KW technetium; tissue; cell type; organ.

OS Synthetic.

XX Key Location/Qualifiers

FT Modified-site 1 /label= Pic-Ser

XX WO9513832-A1.

XX 26-MAY-1995.

XX 16-NOV-1994; 94WO-CA000637.

XX 16-NOV-1993; 93US-00152680.

XX (RESO-) RESOLUTION PHARM INC.

XX Pollak A, Dunn-Dufault R;

XX WPI; 1995-200205/26.

XX Compens. for generating metal labelled imaging agents - comprising a
 FT solid support, a linking gp. and a ligand cleavable from the linking gp.
 FT by a metal.
 XX
 XX
 PS Claim 6; Page 21; 31pp; English.
 XX
 XX Peptides AAS76218-9 are examples of peptide-ligands used in a method to
 CC produce a metal-labelled imaging agent. The method comprises attaching a
 CC ligand-target molecule e.g. a peptide to a solid surface via a linking
 CC group e.g. maleimide. The ligand can incorporate a metal atom e.g.
 CC technetium, which cleaves the linker-ligand bond thus releasing a
 CC labelled peptide. The labelled peptide can then be used for imaging in
 CC tissues, cell types or organs
 XX
 XX Sequence 8 AA;
 SQ

Query Match 100.0%; Score 29; DB 2; Length 8;
 Best Local Similarity 100.0%; Pred. No. 1.4e-06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 Db 4 TKPPR 8

RESULT 25
 AAW11058
 ID AAW11058 standard; peptide; 8 AA.
 XX
 AC AAW11058;
 XX
 DT 03-JUN-1997 (first entry)
 XX
 XX Leukocyte-targeted peptide derived radionuclide chelator.
 XX
 DE Leukocyte; target; direct; chelator; radionuclide; radiolabel; isotope;
 XX diagnostic imaging.
 KW
 KW Synthetic.
 OS
 XX Key Location/Qualifiers
 FT Modified-site 1 /note= "Sarcosine-Ser"
 FT Modified-site 2 /note= "protected with Acn group"
 FT
 FT
 FT WO9603427-A1.
 XX
 XX 08-FEB-1996.
 XX
 PD 28-APR-1995; 95WO-CA000249.
 XX
 PF 22-JUL-1994; 94US-00279155.
 XX
 PR (RESO-) RESOLUTION PHARM INC.
 XX
 PA Pollak A, Goodbody A;
 XX
 PI WPI; 1996-116994/12.
 XX
 DR New peptide derived radionuclide chelators and metal complexes - useful
 XX for diagnostic imaging.
 FT
 PT Example 3; Page 14; 30pp; English.
 PS
 XX AAW11058-W1059 are peptide derived radionuclide chelators that are
 CC targeted to leukocytes, via the GTPPR sequence. When the chelators are
 CC coupled to a targeting molecule and labelled with a diagnostically useful
 CC metal, they can be used to detect pathological conditions by diagnostic
 CC imaging. The leukocyte-targeted chelators are useful for the rapid
 CC imaging of sites of local inflammation. Radionuclides used include 99mTc,
 CC 64Cu, 67Cu, 97Ru, 105Rh, 109Pd, 186Re, 188Re, 198Au, 199Au, 203Pb, 212Pb

CC and 212Bi. The coupling of a targeting agent and radionuclide using a
 CC chelating agent is an alternative to the direct labelling of targeting
 CC agents in which radionuclides are typically bound at the more numerous
 CC low-affinity sites, forming unstable complexes. The new conjugates give
 CC better scintigraphic images in rat inflammation studies than known
 CC imaging agents Ga-67, 99mTc-19G, 111In-WBC and 99mTc-Nanocol. They image
 CC more rapidly than the known agents and show superior biodistribution
 XX
 XX Sequence 8 AA;
 SQ

Query Match 100.0%; Score 29; DB 2; Length 8;
 Best Local Similarity 100.0%; Pred. No. 1.4e-06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 Db 4 TKPPR 8

RESULT 26
 AAY49844
 ID AAY49844 standard; peptide; 8 AA.
 XX
 AC AAY49844;
 XX
 DT 20-JAN-2000 (first entry)
 XX
 XX Tuftsin receptor antagonist chelate conjugate #5.
 DE
 DE Tuftsin receptor antagonist; chelate conjugate; radiopharmaceutical;
 XX diagnosis; infection; inflammation; imaging; cancer; tumour.
 KW
 KW Synthetic.
 OS
 XX Key Location/Qualifiers
 FT Modified-site 1 /note= "6-(hydrazino)-3-pyridylcarbonyl"
 FT Modified-site 2 /note= "modification on the COOH side group: -Acp-Thr-Lys
 FT Modified-site 3 /label= Acp
 FT /note= "6-aminocaproic acid"
 FT
 FT WO9951628-A1.
 XX
 XX 14-OCT-1999.
 XX
 PD 29-MAR-1999; 99WO-US006824.
 XX
 PF 03-APR-1998; 98US-0080672P.
 XX
 PR (DUPO) DU PONT PHARM CO.
 XX
 PA Edwards DS, Rajopadhye M;
 XX
 PI WPI; 1999-633729/54.
 XX
 DR New polypeptides and radiopharmaceuticals used for imaging infection,
 XX inflammation and cancer.
 PT
 PT Claim 4; Page 71; 80pp; English.
 PS
 XX The present invention describes polypeptide compounds of formula (I),
 CC capable of direct transformation into a radiopharmaceutical: Ch-Ln-(X1-X2
 CC -X3-X4-X5)d (I), where X1-X5 = amino acids; Ln = a linking group; Ch = a
 CC metal bonding unit; and d is selected from 1, 2 and 3. The
 CC radiopharmaceuticals are useful for the diagnosis of infection,
 CC inflammation and cancer. The radiopharmaceuticals bind in vivo to the
 CC tuftsin receptor on the surface of white cells which accumulate at the
 CC site of infection and inflammation and can then be detected using
 CC radiation detecting probes or by imaging using a planar or ring gamma
 CC camera. The radiopharmaceuticals can also be used in treating cancer. The

CC present sequence represents a specifically claimed tuftsin receptor
 CC antagonist chelate conjugate from the present invention

XX Sequence 8 AA;

Query Match 100.0%; Score 29; DB 2; Length 8;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 4 TKPPR 8

RESULT 27

ABB08448

ID ABB08448 standard; peptide; 8 AA.

XX

AC ABB08448;

XX 01-JUL-2002 (first entry)

XX Tuftsin receptor antagonist (TKPPR) derivative tetramer (BRU-346).

XX Tuftsin; endothelial cell; drug delivery; gene therapy; NP-1;
 KW angiogenesis; tumour cell; cytostatic; antagonist.

XX

OS Synthetic.

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

CC related disorder. They may be used for delivering desired nucleic acids
 CC to endothelial cells, tumour cells or other cells expressing NP-1, for
 CC enhancing endothelial or tumour cell-targeted gene therapy, or gene
 CC therapy targeting angiogenic cells, and for treating a human or animal
 CC with a tumour or angiogenesis-related disease. The current sequence
 CC represents a tuftsin receptor antagonist (TKPPR) derivative tetramer BRU-
 CC 346

XX Sequence 8 AA;

Query Match 100.0%; Score 29; DB 5; Length 8;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 1 TKPPR 5

RESULT 28

AAR85539

ID AAR85539 standard; peptide; 9 AA.

XX

AC AAR85539;

XX 17-APR-1996 (first entry)

XX Metal chelating peptide conjugated onto tuftsin antagonist.

XX chelate; chelating; chelator; diagnostic; imaging; therapy; tuftsin.

XX Synthetic.

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

CC present sequence represents a specifically claimed tuftsin receptor
 CC antagonist chelate conjugate from the present invention

XX Sequence 8 AA;

Query Match 100.0%; Score 29; DB 2; Length 8;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 4 TKPPR 8

RESULT 27

ABB08448

ID ABB08448 standard; peptide; 8 AA.

XX

AC ABB08448;

XX 01-JUL-2002 (first entry)

XX Tuftsin receptor antagonist (TKPPR) derivative tetramer (BRU-346).

XX Tuftsin; endothelial cell; drug delivery; gene therapy; NP-1;
 KW angiogenesis; tumour cell; cytostatic; antagonist.

XX

OS Synthetic.

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

CC related disorder. They may be used for delivering desired nucleic acids
 CC to endothelial cells, tumour cells or other cells expressing NP-1, for
 CC enhancing endothelial or tumour cell-targeted gene therapy, or gene
 CC therapy targeting angiogenic cells, and for treating a human or animal
 CC with a tumour or angiogenesis-related disease. The current sequence
 CC represents a tuftsin receptor antagonist (TKPPR) derivative tetramer BRU-
 CC 346

XX Sequence 8 AA;

Query Match 100.0%; Score 29; DB 5; Length 8;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 1 TKPPR 5

RESULT 28

AAR85539

ID AAR85539 standard; peptide; 9 AA.

XX

AC AAR85539;

XX 17-APR-1996 (first entry)

XX Metal chelating peptide conjugated onto tuftsin antagonist.

XX chelate; chelating; chelator; diagnostic; imaging; therapy; tuftsin.

XX Synthetic.

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

CC present sequence represents a specifically claimed tuftsin receptor
 CC antagonist chelate conjugate from the present invention

XX Sequence 8 AA;

Query Match 100.0%; Score 29; DB 2; Length 8;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
 |||||
 Db 4 TKPPR 8

RESULT 27

ABB08448

ID ABB08448 standard; peptide; 8 AA.

XX

AC ABB08448;

XX 01-JUL-2002 (first entry)

XX Tuftsin receptor antagonist (TKPPR) derivative tetramer (BRU-346).

XX Tuftsin; endothelial cell; drug delivery; gene therapy; NP-1;
 KW angiogenesis; tumour cell; cytostatic; antagonist.

XX

OS Synthetic.

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

XX

CC binding peptide. The molecule can be used to chelate a diagnostically or
 CC therapeutically useful metal such as 99m-Tc, and the targeting moiety
 CC can be used to direct the molecule to a site of interest within the body
 CC for in-vivo diagnostic imaging or for therapy. The present sequence
 CC represents specifically claimed examples of the new peptides, in which
 CC the targeting peptide is a tuftsin antagonist

SQ Sequence 9 AA;

Query Match 100.0%; Score 29; DB 2; Length 9;
 Best Local Similarity 100.0%; Pred. NO. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TKPPR 5
 |||||
 Db 5 TKPPR 9

RESULT 29
 AAR85535
 ID AAR85535 standard; peptide; 9 AA.
 AC AAR85535;
 XX
 DT 17-APR-1996 (first entry)
 DE Metal chelating peptide conjugated onto tuftsin antagonist.
 XX
 KW chelate; chelating; chelator; diagnostic; imaging; therapy; tuftsin.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT Domain 1..4 /label= metal_chelating_moiety
 FT Modified-site 1 /label= OTHER
 FT Modified-site 3 /note= "picolinic acid residue"
 FT Modified-site 5 /note= "Cys(Acm)"
 FT Domain 5..9 /label= tuftsin_antagonist

XX WO9517419-A1.

PN 29-JUN-1995.

XX 22-DEC-1994; 94WO-CA000718.

XX 22-DEC-1993; 93US-00171737.

XX (RESO-) RESOLUTION PHARM INC.

PA Pollak A, Goodbody A;

PI WPI; 1995-240607/31.

DR New peptide analogue metal chelator(s) - used for chelating radioactive
 PT metals for in-vivo diagnostic imaging or for therapy.

XX Claim 11; Page 26; 34pp; English.

CC The patent discloses new metal-chelating peptide analogues of formula
 CC Chel-Xaa-Cys-X, in which: Chel is a chelating moiety which is the residue
 CC of a heterocyclic amino acid analogue such as picolinic acid, dipicolinic
 CC acid, chelidonic acid, 2-carboxypyrazine, 2-carboxypyrimidine, 2-
 CC carboxypyrrrole, 2-quinolinic acid or 2- or 3-isquinolinic acid; Xaa is
 CC an amino acid residue; the Cys residue is optionally protected; and X is
 CC OH, alkoxyl, an amino acid residue or a targeting moiety (TM). Preferably
 CC Xaa is Gly or Ser; X is Gly-OH or Gly-TM; and TM is a cell receptor
 CC binding peptide. The molecule can be used to chelate a diagnostically or
 CC therapeutically useful metal such as 99m-Tc, and the targeting moiety
 CC can be used to direct the molecule to a site of interest within the body

CC for in-vivo diagnostic imaging or for therapy. The present sequence
 CC represents a specifically claimed example of the new peptides, in which
 CC the targeting peptide is a tuftsin antagonist

SQ Sequence 9 AA;

Query Match 100.0%; Score 29; DB 2; Length 9;
 Best Local Similarity 100.0%; Pred. NO. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TKPPR 5
 |||||
 Db 5 TKPPR 9

RESULT 30
 AAR88735
 ID AAR88735 standard; peptide; 9 AA.
 XX
 AC AAR88735;
 XX
 DT 10-APR-1996 (first entry)
 DE Tuftsin antagonist peptide-metal chelator conjugate.
 XX
 KW Peptide-chelator conjugate; metal chelator; diagnostic imaging;
 XX inflammation; radionuclide; tuftsin; analogue; antagonist.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT Region 1..3 /label= chelator
 FT Misc-difference 1 /note= "pref. Chelates a radionuclide"
 FT Modified-site 3 /label= OTHER
 FT Modified-site 3 /note= "picolinic acid or N',N'-dimethylglycine"
 FT Modified-site 3 /label= OTHER
 FT Region 4 /note= "Cys(Acm)"
 FT Region 5..9 /label= linking_group
 FT Peptide /label= tuftsin_antagonist

XX WO9522996-A2.

PN 31-AUG-1995.

XX 24-FEB-1995; 95WO-CA000106.

XX 25-FEB-1994; 94US-00202178.

XX (RESO-) RESOLUTION PHARM INC.

PA Goodbody A, Pollak A;

PI WPI; 1995-311386/40.

DR New peptide-chelator conjugate and complex with traceable metal - used to
 PT image sites of inflammation in vivo without significant accumulation on
 PT the gastrointestinal tract.

XX Claim 16; Page 21; 23pp; English.

CC The present sequence is that of a specifically claimed peptide- chelator
 CC conjugate in which a tuftsin antagonist peptide is coupled to a metal
 CC chelator, via a linking group. The chelator serves as a labelling site
 CC for radionuclide metals such as technetium-99m. The tuftsin antagonist
 CC targets the conjugate to macrophages and neutrophils at sites of
 CC inflammation without significant accumulation in the gastrointestinal
 CC tract (unlike the native tuftsin tetrapeptide). The conjugate is thus
 CC useful for diagnostic imaging of inflammation sites, providing an

```
CC improved target to background ratio
XX
SQ Sequence 9 AA;
Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. NO. 1.4e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
Db 5 TKPPR 9
RESULT 31
AAR88738
ID AAR88738 standard; peptide; 9 AA.
XX
AC AAR88738;
XX
DT 10-APR-1996 (first entry)
XX
DE Tuftsin antagonist peptide-metal chelator conjugate.
XX
KW Peptide-chelator conjugate; metal chelator; diagnostic imaging;
inflammation; radionuclide; tuftsin; analogue; antagonist.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Region 1..3 /label= chelator
FT /note= "pref. Chelates a radionuclide"
FT Modified-site 1 /label= OTHER
FT /note= "N',N'-dimethylglycine"
FT Modified-site 3 /label= OTHER
FT /note= "Cys(Acm)"
FT Modified-site 4 /label= bala
FT /note= "linking group"
FT Peptide 5..9 /label= tuftsin_antagonist
XX
PN WO9522996-A2.
XX
PD 31-AUG-1995.
XX
PF 24-FEB-1995; 95WO-CA000106.
XX
PR 25-FEB-1994; 94US-00202178.
XX
PA (RESO-) RESOLUTION PHARM INC.
XX
PI Goodbody A, Pollak A;
XX
DR WPI; 1995-311386/40.
XX
PT New peptide-chelator conjugate and complex with traceable metal - used to
image sites of inflammation in vivo without significant accumulation on
the gastrointestinal tract.
XX
PS Claim 16; Page 21; 23pp; English.
XX
CC The present sequence is that of a specifically claimed peptide- chelator
conjugate in which a tuftsin antagonist peptide is coupled to a metal
chelator via a linking group. The chelator serves as a labelling site
for radionuclide metals such as technetium-99m. The tuftsin antagonist
targets the conjugate to macrophages and neutrophils at sites of
inflammation without significant accumulation in the gastrointestinal
tract (unlike the native tuftsin tetrapeptide). The conjugate is thus
useful for diagnostic imaging of inflammation sites, providing an
improved target to background ratio
CC
```

```
XX
SQ Sequence 9 AA;
Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. NO. 1.4e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
Db 5 TKPPR 9
RESULT 32
AAR88741
ID AAR88741 standard; peptide; 9 AA.
XX
AC AAR88741;
XX
DT 10-APR-1996 (first entry)
XX
DE Tuftsin antagonist peptide-metal chelator conjugate.
XX
KW Peptide-chelator conjugate; metal chelator; diagnostic imaging;
inflammation; radionuclide; tuftsin; analogue; antagonist.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Peptide 1..5 /label= tuftsin_antagonist
FT Region 6..7 /label= linking_group
FT Modified-site 6 /label= bala
FT Modified-site 7 /label= bala
FT Modified-site 8 /label= OTHER
FT /note= "the side-chain (epsilon) amino group of Lys at
position 8 forms a peptide bond with the C-terminus of
the tetrapeptide N',N'-dimethylglycyl-Ser-Cys(Acm)-Gly-;
this sidechain group consists of a tripeptide metal
chelator and a linking residue (Gly)"
XX
PN WO9522996-A2.
XX
PD 31-AUG-1995.
XX
PF 24-FEB-1995; 95WO-CA000106.
XX
PR 25-FEB-1994; 94US-00202178.
XX
PA (RESO-) RESOLUTION PHARM INC.
XX
PI Goodbody A, Pollak A;
XX
DR WPI; 1995-311386/40.
XX
PT New peptide-chelator conjugate and complex with traceable metal - used to
image sites of inflammation in vivo without significant accumulation on
the gastrointestinal tract.
XX
PS Claim 19; Page 21; 23pp; English.
XX
CC The present sequence is that of a specifically claimed peptide- chelator
conjugate in which a tuftsin antagonist peptide is coupled to a metal
chelator via a linking group. The chelator serves as a labelling site
for radionuclide metals such as technetium-99m. The tuftsin antagonist
targets the conjugate to macrophages and neutrophils at sites of
inflammation without significant accumulation in the gastrointestinal
tract (unlike the native tuftsin tetrapeptide). The conjugate is thus
useful for diagnostic imaging of inflammation sites, providing an
improved target to background ratio
CC
```

```

XX SQ Sequence 9 AA;
Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.4e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 1 TKPPR 5

RESULT 33
AAR76219
ID AAR76219 standard; peptide; 9 AA.
AC AAR76219;
XX 12-JAN-1996 (first entry)
DT
XX Peptide-ligand conjugate #2.
DE
XX Peptide-ligand; metal-labelled imaging agent; maleimide; metal atom;
KW technetium; tissue; cell type; organ.
XX Synthetic.
OS
XX Key Location/Qualifiers
FH Modified-site 1 /label= N,N'-dimethyl-Gly
FT Modified-site 3
FT Modified-site 3
FT
XX WO9513832-A1.
PN
XX 26-MAY-1995.
PD
XX 16-NOV-1994; 94WO-CA000637.
PF
XX 16-NOV-1993; 93US-00152680.
PR
XX (RESO-) RESOLUTION PHARM INC.
PA Pollak A, Dunn-Dufault R;
PI WPI; 1995-200205/26.
DR
XX Compsns. for generating metal labelled imaging agents - comprising a
PT solid support, a linking gp. and a ligand cleavable from the linking gp.
PT by a metal.
PT
XX Claim 6; Page 21; 31pp; English.
PS
XX Peptides AAR76218-9 are examples of peptide-ligands used in a method to
CC produce a metal-labelled imaging agent. The method comprises attaching a
CC ligand-target molecule e.g. a peptide to a solid surface via a linking
CC group e.g. maleimide. The ligand can incorporate a metal atom e.g
CC technetium, which cleaves the linker-ligand bond thus releasing a
CC labelled peptide. The labelled peptide can then be used for imaging in
CC tissues, cell types or organs
XX
XX Sequence 9 AA;
Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.4e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 5 TKPPR 9

RESULT 34
AAW11055
ID AAW11055 standard; peptide; 9 AA.

```

```

XX AAW11055;
XX 03-JUN-1997 (first entry)
DT
XX Leukocyte-targeted peptide derived radionuclide chelator.
DE
XX Leukocyte; target; direct; chelator; radionuclide; radiolabel; isotope;
KW diagnostic imaging.
XX Synthetic.
XX Key Location/Qualifiers
FH Modified-site 1 /note= "N, N-dimethyl-Gly"
FT Modified-site 3
FT Modified-site 3
FT /note= "protected with Acn group"
XX WO9603427-A1.
PN
XX 08-FEB-1996.
PD
XX 28-APR-1995; 95WO-CA000249.
PF
XX 22-JUL-1994; 94US-00279155.
PR
XX (RESO-) RESOLUTION PHARM INC.
PA Pollak A, Goodbody A;
PI WPI; 1996-116994/12.
DR
XX New peptide derived radionuclide chelators and metal complexes - useful
PT for diagnostic imaging.
PT
XX Claim 28; Page 22; 30pp; English.
PS
XX AAW11054-W11059 are peptide derived radionuclide chelators that are
CC targeted to leukocytes, via the GKPPR sequence. When the chelators are
CC targeted to a targeting molecule and labelled with a diagnostically useful
CC metal, they can be used to detect pathological conditions by diagnostic
CC imaging. The leukocyte-targeted chelators are useful for the rapid
CC imaging of sites of local inflammation. Radionuclides used include 99mTc,
CC 64Cu, 67Cu, 97Ru, 105Rh, 103Pd, 188Re, 188Re, 198Au, 203Pb, 212Pb,
CC and 212Bi. The coupling of a targeting agent and radionuclide using a
CC chelating agent is an alternative to the direct labelling of targeting
CC agents in which radionuclides are typically bound at the more numerous
CC low-affinity sites, forming unstable complexes. The new conjugates give
CC better scintigraphic images in rat inflammation studies than known
CC imaging agents Ga-67, 99mTc-19G, 111In-WBC and 99mTc-Nanocol. They image
CC more rapidly than the known agents and show superior biodistribution
XX
XX Sequence 9 AA;
Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.4e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5
DB 5 TKPPR 9

RESULT 35
AAW11059
ID AAW11059 standard; peptide; 9 AA.
XX
XX AAW11059;
XX 03-JUN-1997 (first entry)
DT
XX Leukocyte-targeted peptide derived radionuclide chelator.
DE
XX

```


KW Leukocyte; target; direct; chelator; radionuclide; radiolabel; isotope;
 KW diagnostic imaging.
 XX
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 FT Modified-site 1
 FT Modified-site /note= "Sarcosine-Gly"
 FT Modified-site 3
 FT /note= "protected with Acm group"
 FT
 XX WO9603427-A1.
 PN
 XX 08-FEB-1996.
 PD
 XX 28-APR-1995; 95WO-CA000249.
 PF
 XX 22-JUL-1994; 94US-00279155.
 PR
 XX (RESO-) RESOLUTION PHARM INC.
 PA
 XX Pollak A, Goodbody A;
 PI
 XX WPI; 1996-116994/12.
 DR
 XX New peptide derived radionuclide chelators and metal complexes - useful
 PT for diagnostic imaging.
 PT
 XX Example 1; Page 12; 30pp; English.
 PS
 XX AAW11054-W11059 are peptide derived radionuclide chelators that are
 CC targeted to leukocytes, via the GRKPPR sequence. When the chelators are
 CC coupled to a targeting molecule and labelled with a diagnostically useful
 CC metal, they can be used to detect pathological conditions by diagnostic
 CC imaging. The leukocyte-targeted chelators are useful for the rapid
 CC imaging of sites of local inflammation. Radionuclides used include 99mTc,
 CC 64Cu, 67Cu, 97Ru, 105Rh, 109Pd, 186Re, 188Re, 198Au, 203Pb, 212Pb
 CC and 212Bi. The coupling of a targeting agent and radionuclide using a
 CC chelating agent is an alternative to the direct labelling of targeting
 CC agents in which radionuclides are typically bound at the more numerous
 CC low-affinity sites, forming unstable complexes. The new conjugates give
 CC better scintigraphic images in rat inflammation studies than known
 CC imaging agents Ga-67, 99mTc-IgG, 111In-WBC and 99mTc-Nanocol. They image
 CC more rapidly than the known agents and show superior biodistribution
 XX
 SQ Sequence 9 AA;
 Query Match 100.0%; Score 29; DB 2; Length 9;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 Db 5 TKPPR 9
 RESULT 36
 AAW11054
 ID AAW11054 standard; peptide; 9 AA.
 XX
 AC AAW11054;
 XX
 XX 03-JUN-1997 (first entry)
 DT
 XX Leukocyte-targeted peptide derived radionuclide chelator.
 DE
 XX Leukocyte; target; direct; chelator; radionuclide; radiolabel; isotope;
 KW diagnostic imaging.
 KW
 XX Synthetic.
 OS
 XX Key Location/Qualifiers
 FT Modified-site 1
 FT Modified-site /note= "N, N-diethyl-Gly"
 FT Modified-site 3
 FT /note= "protected with Acm group"
 FT
 XX WO9603427-A1.
 PN
 XX 08-FEB-1996.
 PD

FT Modified-site /note= "N, N-dimethyl-Gly"
 FT 3
 FT /note= "protected with Acm group"
 XX
 PN WO9603427-A1.
 XX
 XX 08-FEB-1996.
 PD
 XX 28-APR-1995; 95WO-CA000249.
 PF
 XX 22-JUL-1994; 94US-00279155.
 PR
 XX (RESO-) RESOLUTION PHARM INC.
 PA
 XX Pollak A, Goodbody A;
 PI
 XX WPI; 1996-116994/12.
 DR
 XX New peptide derived radionuclide chelators and metal complexes - useful
 PT for diagnostic imaging.
 PT
 XX Claim 28; Page 22; 30pp; English.
 PS
 XX AAW11054-W11059 are peptide derived radionuclide chelators that are
 CC targeted to leukocytes, via the GRKPPR sequence. When the chelators are
 CC coupled to a targeting molecule and labelled with a diagnostically useful
 CC metal, they can be used to detect pathological conditions by diagnostic
 CC imaging. The leukocyte-targeted chelators are useful for the rapid
 CC imaging of sites of local inflammation. Radionuclides used include 99mTc,
 CC 64Cu, 67Cu, 97Ru, 105Rh, 109Pd, 186Re, 188Re, 198Au, 203Pb, 212Pb
 CC and 212Bi. The coupling of a targeting agent and radionuclide using a
 CC chelating agent is an alternative to the direct labelling of targeting
 CC agents in which radionuclides are typically bound at the more numerous
 CC low-affinity sites, forming unstable complexes. The new conjugates give
 CC better scintigraphic images in rat inflammation studies than known
 CC imaging agents Ga-67, 99mTc-IgG, 111In-WBC and 99mTc-Nanocol. They image
 CC more rapidly than the known agents and show superior biodistribution
 XX
 SQ Sequence 9 AA;
 Query Match 100.0%; Score 29; DB 2; Length 9;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 Db 5 TKPPR 9
 RESULT 37
 AAW11056
 ID AAW11056 standard; peptide; 9 AA.
 XX
 AC AAW11056;
 XX
 XX 03-JUN-1997 (first entry)
 DT
 XX Leukocyte-targeted peptide derived radionuclide chelator.
 DE
 XX Leukocyte; target; direct; chelator; radionuclide; radiolabel; isotope;
 KW diagnostic imaging.
 KW
 XX Synthetic.
 OS
 XX Key Location/Qualifiers
 FT Modified-site 1
 FT Modified-site /note= "N, N-diethyl-Gly"
 FT Modified-site 3
 FT /note= "protected with Acm group"
 FT
 XX WO9603427-A1.
 PN
 XX 08-FEB-1996.
 PD

XX PF 28-APR-1995; 95WO-CA000249.
 XX PR 22-JUL-1994; 94US-00279155.
 XX PA (RESO-) RESOLUTION PHARM INC.
 XX PI Pollak A, Goodbody A;
 XX PS WPI; 1996-116994/12.
 XX DR New peptide derived radionuclide chelators and metal complexes - useful
 XX PT for diagnostic imaging.
 XX PS Claim 28; Page 22; 30pp; English.
 XX CC AAW11054-W11059 are peptide derived radionuclide chelators that are
 CC targeted to leukocytes, via the GTPPR sequence. When the chelators are
 CC coupled to a targeting molecule and labelled with a diagnostically useful
 CC metal, they can be used to detect pathological conditions by diagnostic
 CC imaging. The leukocyte-targeted chelators are useful for the rapid
 CC imaging of sites of local inflammation. Radionuclides used include 99mTc,
 CC 64Cu, 67Cu, 97Ru, 105Rh, 109Pd, 186Re, 188Re, 198Au, 203Pb, 212Pb
 CC and 212Bi. The coupling of a targeting agent and radionuclide using a
 CC chelating agent is an alternative to the direct labelling of targeting
 CC agents in which radionuclides are typically bound at the more numerous
 CC low-affinity sites, forming unstable complexes. The new conjugates give
 CC better scintigraphic images in rat inflammation studies than known
 CC imaging agents Ga-67, 99mTc-IGG, 111In-WBC and 99mTc-Nanocol. They image
 CC more rapidly than the known agents and show superior biodistribution
 XX SQ Sequence 9 AA;
 Query Match 100.0%; Score 29; DB 2; Length 9;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 5 TKPPR 9
 RESULT 38
 AAW11057
 ID AAW11057 standard; peptide; 9 AA.
 AC AAW11057;
 DT 03-JUN-1997 (first entry)
 DE Leukocyte-targeted peptide derived radionuclide chelator.
 KW Leukocyte; target; direct; chelator; radionuclide; radiolabel; isotope;
 KW diagnostic imaging.
 XX Synthetic.
 XX Key Location/Qualifiers
 XX FT Modified-site 1 /note= "N, N-dibenzyl-Gly"
 XX FT Modified-site 3 /note= "protected with acm group"
 XX PN WO9603427-A1.
 XX PD 08-FEB-1996.
 XX PO 28-APR-1995; 95WO-CA000249.
 XX PF 22-JUL-1994; 94US-00279155.
 XX PR (RESO-) RESOLUTION PHARM INC.
 XX PA Pollak A;
 XX PI

PI Pollak A, Goodbody A;
 XX WPI; 1996-116994/12.
 XX PT New peptide derived radionuclide chelators and metal complexes - useful
 XX FT for diagnostic imaging.
 XX PS Claim 28; Page 22; 30pp; English.
 XX CC AAW11054-W11059 are peptide derived radionuclide chelators that are
 CC targeted to leukocytes, via the GTPPR sequence. When the chelators are
 CC coupled to a targeting molecule and labelled with a diagnostically useful
 CC metal, they can be used to detect pathological conditions by diagnostic
 CC imaging. The leukocyte-targeted chelators are useful for the rapid
 CC imaging of sites of local inflammation. Radionuclides used include 99mTc,
 CC 64Cu, 67Cu, 97Ru, 105Rh, 109Pd, 186Re, 188Re, 198Au, 203Pb, 212Pb
 CC and 212Bi. The coupling of a targeting agent and radionuclide using a
 CC chelating agent is an alternative to the direct labelling of targeting
 CC agents in which radionuclides are typically bound at the more numerous
 CC low-affinity sites, forming unstable complexes. The new conjugates give
 CC better scintigraphic images in rat inflammation studies than known
 CC imaging agents Ga-67, 99mTc-IGG, 111In-WBC and 99mTc-Nanocol. They image
 CC more rapidly than the known agents and show superior biodistribution
 XX SQ Sequence 9 AA;
 Query Match 100.0%; Score 29; DB 2; Length 9;
 Best Local Similarity 100.0%; Pred. No. 1.4e+06;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB 5 TKPPR 9
 RESULT 39
 AAW03420
 ID AAW03420 standard; peptide; 9 AA.
 AC AAW03420;
 DT 10-OCT-1997 (first entry)
 DE Peptide useful as diagnostic imaging agent.
 KW chelator; ligand.
 XX Synthetic.
 XX Key Location/Qualifiers
 XX FT Region 1.3 /label= chelator
 XX FT Modified-site 1 /note= "N,N-dimethyl-Gly"
 XX FT Modified-site 3 /label= ligand derivatised residue
 XX FT /note= "a galactosyl residue is attached to the thiol
 XX FT group of this Cys via a cleavable maleimido-containing
 XX FT coupling group"
 XX FT Region 5.9 /label= targeting_molecule
 XX PN WO9638185-A1.
 XX PD 05-DEC-1996.
 XX PO 16-MAY-1996; 96WO-CA000310.
 XX PF 31-MAY-1995; 95US-00454859.
 XX PR (RESO-) RESOLUTION PHARM INC.
 XX PA Pollak A;
 XX PI

XX WPI; 1997-042662/04.
XX
XX New cpd. for use in diagnostic imaging - consists of imaging agent, which
PT is chelator for traceable metal, coupled by metal-cleavable bond to
PT ligand which localises at different site.
XX
XX Claim 10-14; Page 16; 26pp; English.
XX
XX The patent discloses new peptides useful for diagnostic imaging. The
CC peptides comprise (1) an imaging agent (a chelator for a traceable metal)
CC which localises selectively at an in-vivo site of diagnostic interest;
CC (2) a ligand which localises at a point (in vivo) removed from the site
CC of diagnostic interest; and (3) a metal-cleavable bond coupling the
CC chelator of the imaging agent to the ligand, this bond being cleaved upon
CC formation of a coordination complex of the metal and chelator. Preferably
CC the chelator is the sequence N,N-dimethyl- Gly-Ser-Cys; the ligand is a
CC galactosyl residue; the traceable metal is ^{99m}Tc; and a targeting
CC molecule (especially TKPPR) is attached to the chelator. Upon labelling,
CC the ligand is cleaved, leaving the labelled imaging agent free to
CC localise at the site of diagnostic interest unhindered, while the ligand
CC and any unlabelled imaging agent are sequestered to the removed site. By
CC sequestering unlabelled imaging agent, the labelled imaging agent does
CC not compete to occupy the site of interest, resulting in images of
CC enhanced resolution. The present sequence is the preferred molecule used
CC for imaging, in which a galactosyl residue is attached to the thiol group
CC of Cys(3) via a cleavable maleimido coupling group
XX
XX Sequence 9 AA;
SQ

Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.4e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
Db |||||
5 TKPPR 9

RESULT 40
AAAY23752
ID AAAY23752 standard; peptide; 9 AA.
AC AAAY23752;
XX
XX 09-SEP-1999 (first entry)
DE Peptide RP502 used to prepare chirally pure peptides.
XX Chirally pure peptide; metal complex; diagnostic imaging; radioimaging.
XX Synthetic.
XX
FH Key Location/Qualifiers
FT Modified-site 1 /note= "DimethylGly"
FT Modified-site 2 /note= "t-butylGly"
FT Modified-site 3 /note= "Acm protecting group attached"
FT
XX WO9933863-A1.
XX
XX 08-JUL-1999.
XX
XX 23-DEC-1998; 98WO-CA001201.
XX
XX 24-DEC-1997; 97US-00997802.
PR 30-DEC-1997; 97CA-0226226.
XX
XX (RESO-) RESOLUTION PHARM INC.
PA Pollak A, Fauconnier T, Wong E;
XX
PI

XX WPI; 1999-419086/35.
XX
XX Chirally pure peptides for preparation of metal complexes, used for
PT diagnostic imaging.
XX
XX Example 1; Page 18; 35pp; English.
XX
XX The specification describes chirally pure peptides which are used for
CC preparation of metal complexes. When converted to complexes with
CC (radioactive) metals or their oxides or nitrides, the peptides are useful
CC for diagnostic (radio)imaging (by labeling biomolecules). The present
CC peptide is used in the course of the invention
XX
XX Sequence 9 AA;
SQ

Query Match 100.0%; Score 29; DB 2; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.4e+06;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
Db |||||
5 TKPPR 9

RESULT 41
AAR88739
ID AAR88739 standard; peptide; 10 AA.
AC AAR88739;
XX
XX 10-APR-1996 (first entry)
DE Tuftsin antagonist peptide-metal chelator conjugate.
XX Peptide-chelator conjugate; metal chelator; diagnostic imaging;
KW inflammation; radionuclide; tuftsin; analogue; antagonist.
XX
XX Synthetic.
XX
FH Key Location/Qualifiers
FT Region 1, 3
FT /label= chelator
FT /note= "pref. chelates a radionuclide"
FT Modified-site 1 /label= OTHER
FT Modified-site 3 /note= "N',N'-dimethylglycine"
FT /label= OTHER
FT /note= "Cys(Acm)"
FT Region 4, 5
FT /label= linking_group
FT Modified-site 4 /label= bala
FT Modified-site 5 /label= bala
FT Peptide 6, 10
FT /label= tuftsin_antagonist
XX
XX WO9522996-A2.
XX
XX 31-AUG-1995.
XX
XX 24-FEB-1995; 95WO-CA000106.
XX
XX 25-FEB-1994; 94US-00202178.
PR (RESO-) RESOLUTION PHARM INC.
XX
XX Goodbody A, Pollak A;
XX
XX WPI; 1995-311386/40.
XX
XX

PT New peptide-chelator conjugate and complex with traceable metal - used to
XX image sites of inflammation in vivo without significant accumulation on
XX the gastrointestinal tract.
XX
XX Claim 16; Page 21; 23pp; English.
XX
XX The present sequence is that of a specifically claimed peptide- chelator
XX conjugate in which a tuftsin antagonist peptide is coupled to a metal
XX chelator, via a linking group. The chelator serves as a labelling site
XX for radionuclide metals such as technetium-99m. The tuftsin antagonist
XX targets the conjugate to macrophages and neutrophils at sites of
XX inflammation without significant accumulation in the gastrointestinal
XX tract (unlike the native tuftsin tetrapeptide). The conjugate is thus
XX useful for diagnostic imaging of inflammation sites, providing an
XX improved target to background ratio
XX
SQ Sequence 10 AA;
Query Match 100.0%; Score 29; DB 2; Length 10;
Best Local Similarity 100.0%; Pred. No. 72;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 1 TKPPR 5
Db 6 TKPPR 10
RESULT 42
ADE81165
ID ADE81165 standard; peptide; 10 AA.
XX
XX ADE81165;
XX
XX 29-JAN-2004 (first entry)
XX
XX Tuftsin metalloptide analogue #48.
XX
XX immunostimulant; immunosuppressive; tuftsin receptor binder;
XX tuftsin receptor; tuftsin receptor analogue; imaging; infection;
XX inflammation; immunostimulatory; immune system disorder; analgesic;
XX CNS condition; whole body imaging; radiotherapy;
XX tuftsin metalloptide analogue.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX Modified-site 5..6 /note= "Residues joined by NH-(CH2)6-CO"
XX Misc-difference 7 /note= "D-form residue"
XX Misc-difference 9 /note= "D-form residue"
XX
XX US2003059422-A1.
XX
XX 27-MAR-2003.
XX
XX 30-AUG-1999; 99US-00387715.
XX
XX 07-JUN-1995; 95US-00476652.
XX 05-JUN-1996; 96US-00660697.
XX 18-MAR-1998; 98US-0078373P.
XX 14-DEC-1998; 98US-0112235P.
XX 18-MAR-1999; 99WO-US005693.
XX (SHAR/) SHARMA S. D.
XX
XX Sharma SD;
XX
XX WPI; 2003-596563/56.
XX
XX Tuftsin metalloptide, useful e.g. for imaging site of infection or
XX inflammation, comprises metal ion-binding backbone including at least two

PT contiguous amino acids for complexing with metal ion.
XX
XX Claim 17; Page 11; 12pp; English.
XX
XX The invention describes a peptide or its salt comprising a metal ion-
XX binding backbone including at least two contiguous amino acids for
XX complexing with the metal ion. The peptide is specific for the tuftsin
XX receptor on complexing the metal ion-binding backbone with the metal ion.
XX The peptide is useful for imaging a site of infection or inflammation;
XX for causing an immunostimulatory response in mammals; for treating immune
XX system disorders; in biological, pharmaceutical and radiopharmaceutical
XX applications; as an analgesic in the treatment of CNS conditions; and in
XX whole body imaging and radiotherapy. The peptide complexed with the metal
XX ion is resistant to enzymatic degradation. The affinity of the peptide
XX for the tuftsin receptor is higher when the metal ion-binding backbone is
XX complexed with the metal ion than that when the metal ion-binding
XX backbone is not complexed with the metal ion. The Tc-labeled Thr-D-Lys-
XX Gly-D-Cys-Arg is the most potent existing tuftsin molecule. The peptide-
XX metal ion complexes have a higher level of stability, and are less
XX susceptible to proteolysis than the uncomplexed or existing peptides. The
XX peptide analogue is not conformationally restricted in the absence of a
XX metal ion but has high potency and concomitant conformational restriction
XX on complexation with a metal ion. The metal complexation in the peptide
XX causes specific regional conformational restrictions in the peptide, so
XX that the peptide conformation at the metal binding site is
XX conformationally fixed on metal complexation. The complexation of the
XX peptide to a metal ion alters the in vivo distribution profile, rate and
XX mode of clearance from the body, bioavailability and pharmacokinetics in
XX mammals. The peptide metal ion complex stimulates polymorphonuclear
XX granulocytes, monocytes and macrophages towards phagocytosis, produces
XX higher titer antibodies, and can transit the gut-blood barrier without
XX significant enzymatic or peptidase degradation. This is the amino acid
XX sequence of a tuftsin metalloptide analogue of the invention.
XX
SQ Sequence 10 AA;
Query Match 100.0%; Score 29; DB 7; Length 10;
Best Local Similarity 100.0%; Pred. No. 72;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 1 TKPPR 5
Db 1 TKPPR 5
RESULT 43
ADE81166
ID ADE81166 standard; peptide; 10 AA.
XX
XX ADE81166;
XX
XX 29-JAN-2004 (first entry)
XX
XX Tuftsin metalloptide analogue #49.
XX
XX immunostimulant; immunosuppressive; tuftsin receptor binder;
XX tuftsin receptor; tuftsin receptor analogue; imaging; infection;
XX inflammation; immunostimulatory; immune system disorder; analgesic;
XX CNS condition; whole body imaging; radiotherapy;
XX tuftsin metalloptide analogue.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
XX Misc-difference 2 /note= "D-form residue"
XX Misc-difference 4 /note= "D-form residue"
XX Modified-site 5..6 /note= "Residues joined by NH-(CH2)6-CO"
XX
XX US2003059422-A1.

PD 27-MAR-2003.
 XX 30-AUG-1999; 99US-00387715.
 XX
 XX 07-JUN-1995; 95US-00478652.
 XX 05-JUN-1996; 96US-00660697.
 XX 18-MAR-1998; 98US-0078373P.
 XX 14-DEC-1998; 98US-0112235P.
 XX 18-MAR-1999; 99WO-US005693.
 XX (SHAR/) SHARMA S D.
 XX Sharma SD;
 XX WPI; 2003-596563/56.
 XX
 XX Tuftsin metalloproteinase, useful e.g. for imaging site of infection or
 XX inflammation, comprises metal ion-binding backbone including at least two
 XX contiguous amino acids for complexing with metal ion.
 XX
 XX Claim 17; Page 11; 12pp; English.
 XX
 XX The invention describes a peptide or its salt comprising a metal ion-
 XX binding backbone including at least two contiguous amino acids for
 XX complexing with the metal ion. The peptide is specific for the tuftsin
 XX receptor on complexing the metal ion-binding backbone with the metal ion.
 XX The peptide is useful for imaging a site of infection or inflammation;
 XX for causing an immunostimulatory response in mammals; for treating immune
 XX system disorders; in biological, pharmaceutical and radiopharmaceutical
 XX applications; as an analgesic in the treatment of CNS conditions; and in
 XX whole body imaging and radiotherapy. The peptide complexed with the metal
 XX ion is resistant to enzymatic degradation. The affinity of the peptide
 XX for the tuftsin receptor is higher when the metal ion-binding backbone is
 XX complexed with the metal ion than that when the metal ion-binding
 XX backbone is not complexed with the metal ion. The Tc-labeled Thr-D-Lys-
 XX Gly-D-Cys-Arg is the most potent existing tuftsin molecule. The peptide-
 XX metal ion complexes have a higher level of stability, and are less
 XX susceptible to proteolysis than the uncomplexed or existing peptides. The
 XX peptide analogue is not conformationally restricted in the absence of a
 XX metal ion but has high potency and concomitant conformational restriction
 XX on complexation with a metal ion. The metal complexation in the peptide
 XX causes specific regional conformational restrictions in the peptide, so
 XX that the peptide conformation at the metal binding site is
 XX conformationally fixed on metal complexation. The complexation of the
 XX peptide to a metal ion alters the in vivo distribution profile, rate and
 XX mode of clearance from the body, bioavailability and pharmacokinetics in
 XX mammals. The peptide metal ion complex stimulates polymorphonuclear
 XX granulocytes, monocytes and macrophages towards phagocytosis, produces
 XX higher titer antibodies, and can transit the gut-blood barrier without
 XX significant enzymatic or peptidase degradation. This is the amino acid
 XX sequence of a tuftsin metalloproteinase analogue of the invention.
 XX
 XX SQ Sequence 10 AA;
 Query Match 100.0%; Score 29; DB 7; Length 10;
 Best Local Similarity 100.0%; Pred. No. 72;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB |||||
 6 TKPPR 10
 RESULT 44
 AAR88736
 ID AAR88736 standard; peptide; 11 AA.
 XX
 XX AAR88736;
 XX
 XX 10-APR-1996 (first entry)
 DT
 XX Tuftsin antagonist peptide-metal chelator conjugate.
 DE
 XX

KW Peptide-chelator conjugate; metal chelator; diagnostic imaging;
 KW inflammation; radionuclide; tuftsin; analogue; antagonist.
 XX Synthetic.
 OS
 XX Key Location/Qualifiers
 XX Region 1. .3
 FT /label= chelator
 FT /note= "pref. chelates a radionuclide"
 FT Misc-difference 1
 FT /label= OTHER
 FT /note= "picolinic acid or benzoyl-mercaptopoacetic acid"
 FT Modified-site 3
 FT /label= OTHER
 FT /note= "Cys (Acm) "
 FT Region 4. .6
 FT /label= linking_group
 FT Peptide 7. .11
 FT /label= tuftsin_antagonist
 XX WO9522996-A2.
 PN
 XX 31-AUG-1995.
 XX
 XX 24-FEB-1995; 95WO-CA000106.
 XX
 XX 25-FEB-1994; 94US-00202178.
 XX (RESO-) RESOLUTION PHARM INC.
 XX
 XX Goodbody A, Pollak A;
 XX WPI; 1995-311386/40.
 XX
 XX New peptide-chelator conjugate and complex with traceable metal - used to
 XX image sites of inflammation in vivo without significant accumulation on
 XX the gastrointestinal tract.
 XX
 XX Claim 16; Page 21; 23pp; English.
 XX
 XX The present sequence is that of a specifically claimed peptide-chelator
 XX conjugate in which a tuftsin antagonist peptide is coupled to a metal
 XX chelator, via a linking group. The chelator serves as a labelling site
 XX for radionuclide metals such as technetium-99m. The tuftsin antagonist
 XX targets the conjugate to macrophages and neutrophils at sites of
 XX inflammation without significant accumulation in the gastrointestinal
 XX tract (unlike the native tuftsin tetrapeptide). The conjugate is thus
 XX useful for diagnostic imaging of inflammation sites, providing an
 XX improved target to background ratio
 XX
 XX SQ Sequence 11 AA;
 Query Match 100.0%; Score 29; DB 2; Length 11;
 Best Local Similarity 100.0%; Pred. No. 79;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TKPPR 5
 DB |||||
 7 TKPPR 11
 RESULT 45
 ADD10691
 ID ADD10691 standard; peptide; 11 AA.
 XX
 XX ADD10691;
 XX
 XX 01-JAN-2004 (first entry)
 DT
 XX Tuftsin analogue peptide tetramer.
 DE
 XX
 XX Phagocytosis; tuftsin; endothelial cell; inflammation; cytostatic;
 KW antiangiogenic; NP-1; ultrasound contrast agent; tumour; angiogenesis;
 XX

visualisation therapy; radiotherapy.
Synthetic.

Key Modified-site Location/Qualifiers
6
/label= OTHER
/notes "Lys is covalently linked to a further TKPPRKRPPKT
peptide whose Lys (6) is linked to Gly-TTDA (4,7,10-
Trioxal,13-Tridecanediamine)-oregon green"

US2002147136-A1.
10-OCT-2002.
04-JUN-2001; 2001US-00871974.
02-JUN-2000; 2000US-00585364.
(VWRO//) VON WRONSKI M A.
(MARI//) MARINELLI E R.
(NUNN//) NUNN A D.
(PILL//) PILLAI R.
(RAMA//) RAMALINGAM K.
(TWEED//) TWEEDLE M F.
(LIND//) LINDER K.
(NANJ//) NANJAPPAN P.
(RAJU//) RAJU N.

Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;
Tweedle MF, Linder K, Nanjappan P, Raju N;
WPI; 2003-800817/75.

Composition used in targeting endothelial cells e.g. tumor cells
comprises compounds containing monomers, multimers or polymers of L-
arginine-L-threonyl-L-lysyl-L-prolyl-L-prolyl.

Example 29; Page 66; 85pp; English.

The invention relates to a composition (A1) comprising compounds
containing monomers, multimers or polymers of TKPPR (ADD10684).
Composition (A1) comprises a compound of formula A-L-B 1, where A is the
TKPPR peptide, L is a linker moiety (of formula given in the
specification), and B is a substrate (or a phospholipid group,
derivatisable bead attached to a fluorescent or radioactive marker,
bioactive agent, delivery vehicle for genetic material, drug or
therapeutic, or chelating group (preferably N 4, S 4, N 3 S, N 2 S 2 or
NS 3) comprising oxa-PnAO complexed with 99m Tc). The compound
specifically binds to NP-1 (Vascular endothelial growth factor binding
receptor transmembrane glycoprotein) or cells that express NP-1 with
avidity of at least that of TKPPR. Also included are an ultrasound
contrast agent (C1) comprising a suspension of gas filled microbubbles
comprising the TKPPR compound, an ultrasound contrast agent (C2)
comprising a suspension of gas filled microballoons comprising the TKPPR
compound, preparation of the TKPPR compound (which comprises conjugating
the monomer, multimer or polymer of TKPPR or its analogue with a linker
to obtain a compound of formula A-L, forming a covalent or non-covalent
bond between A-L and the substrate B 1 or forming a covalent bond between
B 1 and the linker to form a conjugate B-L followed by conjugation with
the monomer), and a kit for preparing a radiopharmaceutical comprising
the compound. The compound used for targeting endothelial cells, tumour
cells or other cells which express NP-1, for inhibiting angiogenesis, for
ultrasound imaging, staging a tumour, screening at least one targeted
ultrasound contrast agent for the ability to target endothelial cells,
tumour cells or other cells which express NP-1, for the therapeutic
delivery in vivo of a bioactive agent and for delivering desired nucleic
acids to endothelial cells, tumour cells or other cells which express NP-
1. The composition is also useful for visualisation therapy or
radiotherapy of endothelial cells. The present sequence is a TKPPR
tetramer peptide.

Sequence 11 AA;

Query Match 100.0%; Score 29; DB 7; Length 11;
Best Local Similarity 100.0%; Pred. No; 79;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TKPPR 5
|
|
|
|
|
Db 1 TKPPR 5

RESULT 46
ADD10690
ID ADD10690 standard; peptide; 11 AA.
XX
XX AC ADD10690;
XX
XX DT 01-JAN-2004 (first entry)
XX
XX DE Tuftsin analogue peptide dimer.
XX
XX XX Phagocytosis; tuftsin; endothelial cell; inflammation; cytostatic;
XX KW antiangiogenic; NP-1; ultrasound contrast agent; tumour; angiogenesis;
XX KW visualisation therapy; radiotherapy.
XX
XX OS Synthetic.
XX
XX XX Key Location/Qualifiers
XX FT Modified-site 6
XX FT /label= OTHER
XX FT /notes "Lys is covalently linked to Gly-TTDA (4,7,10-
XX FT Trioxal,13-Tridecanediamine)"
XX
XX FN US2002147136-A1.
XX
XX PD 10-OCT-2002.
XX
XX PF 04-JUN-2001; 2001US-00871974.
XX
XX PR 02-JUN-2000; 2000US-00585364.
XX
XX PA (VWRO//) VON WRONSKI M A.
XX PA (MARI//) MARINELLI E R.
XX PA (NUNN//) NUNN A D.
XX PA (PILL//) PILLAI R.
XX PA (RAMA//) RAMALINGAM K.
XX PA (TWEED//) TWEEDLE M F.
XX PA (LIND//) LINDER K.
XX PA (NANJ//) NANJAPPAN P.
XX PA (RAJU//) RAJU N.

CC contrast agent (c1) comprising a suspension of gas filled microbubbles
 CC comprising the TKPPR compound, an ultrasound contrast agent (c2)
 CC comprising a suspension of gas filled microballoons comprising the TKPPR
 CC compound, preparation of the TKPPR compound (which comprises conjugating
 CC the monomer, multimer or polymer of TKPPR or its analogue with a linker
 CC to obtain a compound of formula A-L, forming a covalent or non-covalent
 CC bond between A-L and the substrate B 1 or forming a covalent bond between
 CC B 1 and the linker to form a conjugate B-L followed by conjugation with
 CC the monomer), and a kit for preparing a radiopharmaceutical comprising
 CC the compound, and a kit for preparing a radiopharmaceutical comprising
 CC cells or other cells which express NP-1, for inhibiting angiogenesis, for
 CC ultrasound imaging, staging a tumour, screening at least one targeted
 CC ultrasound contrast agent for the ability to target endothelial cells,
 CC tumour cells or other cells which express NP-1, for the therapeutic
 CC delivery in vivo of a bioactive agent and for delivering desired nucleic
 CC acids to endothelial cells, tumour cells or other cells which express NP-
 CC 1. The composition is also useful for visualisation therapy or
 CC radiotherapy of endothelial cells. The present sequence is a TKPPR dimer
 CC peptide.
 XX
 XX

SQ Sequence 11 AA;

Query Match 100.0%; Score 29; DB 7; Length 11;
 Best Local Similarity 100.0%; Pred. No. 79;
 Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5

Db 1 TKPPR 5

RESULT 47

ABB08446
 ID ABB08446 standard; peptide; 12 AA.

XX ABB08446;

XX 01-JUL-2002 (first entry)

XX Tuftsin receptor antagonist (TKPPR) derivative peptide 3.

XX Tuftsin, endothelial cell; drug delivery; gene therapy; NP-1;
 KW angiogenesis; tumour cell; cytostatic; antagonist.

XX Synthetic.

XX Key Location/Qualifiers

FT Modified-site 1 /note= "residue modified by the following; F-108,
 FT OCH2CONH"

FT Modified-site 2 /note= "residue modified by (tBu)"

FT Modified-site 3 /note= "residue modified by (Wtt)"

FT Modified-site 6 /note= "residue modified by (Wtt)"

FT Modified-site 7 /note= "residue modified by (Pmc)-OtBu"

FT Modified-site 8 /note= "residue modified by the following; F-108,
 FT OCH2CONH"

FT Modified-site 9 /note= "residue modified by (tBu)"

FT Modified-site 12 /note= "residue modified by (Wtt)"

FT Modified-site 12 /note= "residue modified by (Pmc)-OtBu"

XX WO200191805-A2.

PN 06-DEC-2001.

XX 04-JUN-2001; 2001WO-US018053.

XX 02-JUN-2000; 2000US-00595364.

PR (BRAC) BRACCO RES USA.

XX

PA (BRAC) BRACCO RES USA.

XX

XX Von Wronski MA, Marinelli ER, Nunn AD, Pillai R, Ramalingam K;

PI

PI Tweedle MF, Linder K, Nanjappan P, Raju N;

XX

XX WPI; 2002-195523/25.

DR

XX Composition for use in targeting endothelial cells, tumor cells or other

PT

PT cells which express NP-1 comprises a compound containing a polypeptide,

PT

PT linker and substrate.

XX

XX Example 25; Page 95; 146pp; English.

XX

XX The invention relates to a composition for use in targeting endothelial
 CC cells, tumour cells, or other cells which express NP-1. The activity of
 CC compositions of the invention may be described as cytostatic. Compounds
 CC of the invention are useful in pharmaceutical compositions for inhibiting
 CC angiogenesis, for imaging and targeting an angiogenic site, endothelial
 CC cells, tumour cells or other cells that express NP-1 in a human or
 CC animal. They may also be used as ultrasound contrast agents, for staging
 CC a tumour in a human or animal, for screening for the ability of an agent
 CC to target endothelial cells, tumour cells or other cells that express NP-
 CC 1. They may be used for therapeutic delivery in vivo of a bioactive agent
 CC or for treating an individual exhibiting effects of an angiogenesis or a
 CC related disorder. They may be used for delivering desired nucleic acids
 CC to endothelial cells, tumour cells or other cells expressing NP-1, for
 CC enhancing endothelial or tumour cell-targeted gene therapy, or gene
 CC therapy targeting angiogenic cells, and for treating a human or animal
 CC with a tumour or angiogenesis-related disease. The current sequence
 CC represents a tuftsin receptor antagonist (TKPPR) derivative of the
 CC invention
 XX
 XX

SQ Sequence 12 AA;

Query Match 100.0%; Score 29; DB 5; Length 12;

Best Local Similarity 100.0%; Pred. No. 85;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TKPPR 5

Db 2 TKPPR 6

RESULT 48

ABB08449

ID ABB08449 standard; peptide; 20 AA.

XX

AC ABB08449;

XX

XX 01-JUL-2002 (first entry)

DT

XX 99mTc radiopharmaceutical.

DE

XX Tuftsin, endothelial cell; drug delivery; gene therapy; NP-1;

XX

XX angiogenesis; tumour cell; cytostatic; antagonist; radiopharmaceutical.

XX

XX Synthetic.

OS

XX Key Location/Qualifiers

FT Modified-site 1 /note= "residue modified by the addition of 99mTc-Oxa

FT FNAO"

FT

XX WO200191805-A2.

XX

XX 06-DEC-2001.

PD

XX 04-JUN-2001; 2001WO-US018053.

XX

XX 02-JUN-2000; 2000US-00585364.

XX

PA (BRAC) BRACCO RES USA.


```

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder.
XX
XX Homo sapiens.
OS
XX WO200175067-A2.
FN
XX 11-OCT-2001.
PD
XX 30-MAR-2001; 2001WO-US008631.
PF
XX 31-MAR-2000; 2000US-00540217.
PR
XX 23-AUG-2000; 2000US-00649167.
PR
XX (HYSE-) HYSEQ INC.
PA
XX Drmanac RT, Liu C, Tang YT;
PI
XX WPI: 2001-639362/73.
PI
XX N-PSDB; AAS71194.
PI
XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity.
PT
XX Claim 20; SEQ ID NO 37366; 103pp; English.
PS
XX The invention relates to isolated polynucleotide (I) and polypeptide (II)
CC sequences. (I) is useful as hybridisation probes, polymerase chain
CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping.
CC and in recombinant production of (II). The polynucleotides are also used
CC in diagnostics as expressed sequence tags for identifying expressed
CC genes. (I) is useful in gene therapy techniques to restore normal
CC activity of (II) or to treat disease states involving (II). (II) is
CC useful for generating antibodies against it, detecting or quantitating a
CC polypeptide in tissue, as molecular weight markers and as a food
CC supplement. (II) and its binding partners are useful in medical imaging
CC of sites expressing (II). (I) and (II) are useful for treating disorders
CC involving aberrant protein expression or biological activity. The
CC polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG0010-ABG30377 represent novel human diagnostic
CC amino acid sequences of the invention. Note: The sequence data for this
CC patent did not appear in the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 65 AA;
Query Match 100.0%; Score 29; DB 4; Length 65;
Best Local Similarity 100.0%; Pred. No. 3.9e+02;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TKPPR 5
DB 27 TKPPR 31
Search completed: March 3, 2004, 12:17:18
Job time : 60 secs

```